Core obesity model to assess the cost-effectiveness of weight management interventions

Maria De Francesco¹, Sandra Lopes², Henrik H Meincke², Gabriela Vega-Hernandez², Mark Lamotte¹, Michael E J Lean³ ¹IQVIA, HEOR/HTA, Zaventem, Belgium; ²Novo Nordisk A/S, Søborg, Denmark; ³Human Nutrition, University of Glasgow, Glasgow, UK

Background

• Elevated body mass index (BMI) increases morbidity and mortality from chronic diseases including type 2 diabetes (T2D), cardiovascular disease (CVD), musculoskeletal disorders, hypertension, gallstones and some cancers.^{1–3}



Objectives

- The objective of this study was to develop a 'core obesity model' to assess long-term complications and compare costs and benefits of different weight management interventions.
- This poster explores the model structure, its inputs and its clinical outcomes, using an illustrative example.

Methods

- A Markov model was developed (Excel[™] 2013; MicroSoft, Redmond, WA, USA) to estimate the costs and effects of a cohort with obesity over a life-time time horizon.
- The model cycle length is 3 months for the first year, allowing for a treatment 'stopping rule' at 12 weeks, and annually thereafter.
- The model is able to compute results for time horizons ranging from 1 to 40 years.

Health states

- Health states are additive and reflective of complications that:
- » are highly related to obesity according to the World Health Organization⁴ (Figure 1);
- » respond to weight loss;
- » have substantial effects on costs, quality of life (QoL) and/or life expectancy.
- A graphical representation of the model structure is presented in Figure 1.

Mortality



Treatment effect on HbA_{1c} levels is only considered in populations with T2D. Obstructive sleep apnoea, bariatric surgery and knee replacement may occur in any health state and any cycle of the model.

ACS, acute coronary syndrome; BMI, body mass index; HbA_{1c}, glycated haemoglobin, HDL, high-density lipoprotein; OSA, obstructive sleep apnoea; SBP, systolic blood pressure; T2D, type 2 diabetes; TIA, transient ischaemic attack.

Table 1: Source of data determining risk equations used in the model.

Outcome	Risk equation		
T2D	<u>QDiabetes</u> ⁵ , Framingham Offspring ⁷		
CVD (primary prevention)	In NGT: <u>QRisk</u> ⁸ , Framingham Heart Study: equation with BMI ⁹ ; In T2D: <u>UKPDS82</u> ¹⁰ , QRisk ⁸ , Swedish NDR ¹¹		
CVD (secondary prevention)	In NGT: <u>Framingham Recurring Coronary Heart Disease (CHD)</u> 12; In T2D: Framingham Recurring Coronary Heart Disease (CHD) ¹² , <u>UKPDS82</u> 10		
Colorectal cancer	NI-AARP cohort ¹³ , Schlesinger <i>et al.</i> 2015 ¹⁴		
Post-menopausal breast cancer	Renehan <i>et al.</i> 2008 ¹⁵ , Ahn <i>et al.</i> 2007 ¹⁶		
Post-menopausal endometrial cancer	Renehan <i>et al.</i> 2008 ¹⁵ , Million Women Study ¹⁷		
OSA	Sleep AHEAD ¹⁸ , Sleep Heart Health Study ¹⁹		
Knee replacement surgery	Wendelboe et al. 2003 ²⁰		
Risk equations used in the illustrative example are coloured. CVD refers to both ACS and stroke.			

BMI, body mass index; CVD, cardiovascular disease; NGT, normal glucose tolerance; OSA, obstructive sleep apnoea; T2D, type 2 diabetes.

- General population age and gender-specific all-cause mortality is included in the model, based on country specific life-tables.
- Both long-term and short-term mortality are considered.

Transitions

- Transitions describe the progression of the cohort between health states.
- A systematic review of the literature in 2017 identified risk equations, which can inform the transition probabilities between health states (Table 1).

Clinical and economic outcomes

- The clinical outcomes of the model include:
- » cumulative incidence of obesity-related complications;
- » QoL;
- » life expectancy;
- » quality-adjusted life-expectancy.
- The economic outcomes of the model include:
- » incremental cost:utility ratio;
- » net monetary benefit
- Example risk probability of developing T2D and QoL as a function of BMI are shown in Figure 2A and 2B, respectively.
- Baseline characteristics of the model starting cohort (European NGT population) were defined at model entry (Table 2).

Table 2: Baseline characteristics of the model starting cohort

Characteristic	Cohort 1	Cohort 2
Age, years	45	-
	70.0	

Figure 2: (A) Risk of progression to T2D as a function of BMI from the QDiabetes⁵ study; (B) Estimated relationship between BMI and EQ-5D score after controlling for confounding factors from Søltoft *et al.* 2009.⁶





BMI, body mass index; EQ-5D, EuroQol 5D health questionnaire; NGT, normal glucose tolerance; T2D, type 2 diabetes. **Figure 3**: Change in the incidence of obesity-associated complications in the illustrative example.





Stroke includes TIA.

ACS, acute coronary syndrome; T2D, type diabetes; TIA, transient ischaemic attack; TKR, total knee replacement.

Discussion/Limitations

- Prediabetes is defined at baseline; however, no risk equations for developing prediabetes were identified.
- » Although its management costs are low, this potentially underestimates future prevalence.
- Some risk equations, for example UKPDS82, do not include BMI as an independent risk factor predicting the risk of CVD.

Terriales, 70	70.0	-
Smokers, %	15.9	-
BMI, kg/m ²	37.0	34.0
SBP, mmHg	124.1	-
TC, mmol/L	5.0	-
HDL-C, mmol/L	1.3	-
TG level ≥1.7 mmol/L, %	36	-

Cohort 2 were assumed to have the same the same baseline characteristics as cohort 1, with the exception of a reduction in BMI to 34 kg/m². BMI, body mass index; HDL-C, high density lipoprotein cholesterol; SBP, systolic blood pressure; TC, total cholesterol; TG, triglyceride.

Results

- Over a 10-year time horizon, the model predicted 5.0% mortality, 5.8% acute coronary syndrome (ACS) and 2.1% stroke (Figure 3).
 - » Knee replacement occurred in 6.3% of the cohort and the proportion alive with T2D was 5.0%.
- Following a reduction of 3 BMI units, mortality dropped to 4.8%, ACS to 5.5% and stroke incidence to 2.0%.
- » The incidence of knee replacement was reduced to 4.4% and the proportion alive with T2D was 3.8%.

an macpenderte list factor predicting the list of ever

- » Other risk equations include BMI but only up to certain BMI levels.
- Finally, this study is limited by lack of prospective long-term data on hard outcomes after intentional weight changes.

Conclusion

- The presented Core Obesity Model is novel in assessing the longterm effects of weight management interventions on such a comprehensive set of obesity medical complications.
- This model could therefore be used to inform cost-effectiveness analyses on treatments for adult patients with obesity.

Novo Nordisk provided full sponsorship for this study.

Michael Lean has received departmental research funding and contributed to Advisory Boards for Novo Nordisk. Mark Lamotte is an employee of IQVIA. Sandra Lopes and Henrik Hjorth Meincke are employees of Novo Nordisk. At the time of model development, Maria De Francesco was an employee of IQVIA and Gabriela Vega-Hernandez was an employee of Novo Nordisk. The authors are grateful to Jamie Cozens of Watermeadow Medical (supported by Novo Nordisk), for writing assistance. Presented at the International Society for Pharmacoeconomics and Outcomes Research (ISPOR) Europe 2018. 10–14 November 2018, Barcelona, Spain. References: (1) Guh *et al. BMC Public Health* 2009;9:88; (2) Bhaskaran *et al. Lancet* 2014;384:755–65; (3) Berrington de Gonzalez *et al. N Engl J Med* 2010;363:2211–19; (4) World Health Organization. WHO Consultation on Obesity. 2000. Available at: http://www.who.int/nutrition/publications/obesity/WHO_TRS_894/en; (5) Hippisley-Cox *et al. BMJ* 2017;359:j5019; (6) Soltoft *et al. Qual Life Res* 2009;18:1293–99; (7) Wilson *et al. Arch Intern Med* 2007;167:1068–74; (8) Hippisley-Cox *et al. BMJ* 2017;357:j2099; (9) D'Agostino *et al. Circulation* 2008;117:743–53; (10) McEwan *et al. Cost Eff Resour Alloc* 2015;13:1; (11) Cederholm *et al. Diab Care* 2008;31:2038–43; (12) D'Agostino *et al. Am J Heart* 2000;139:272–81; (13) Adams *et al. Am J Epidemiol* 2007;166:36–45; (14) Schlesinger *et al. Obes Rev* 2015;16:607–19; (15) Renehan *et al. Lancet* 2008;371:569–78; (16) Ahn *et al. Arch Intern Med* 2007;167:2091–102; (17) Yang *et al. Br J Cancer* 2012;107:169–75; (18) Kuna *et al. Sleep* 2013;36:641–9; (19) Young *et al. Arch Intern Med* 2002;162:893–900; (20) Wendelboe *et al. Am J Prev Med* 2003;25:290–5.