

Identification and expert panel rating of key structural approaches applied in health economic obesity models

Schwander, Bjorn ; Nuijten, Mark ; Hiligsmann, Mickaël ; Queally, Michelle ; Leidl, Reiner ; Joore, Manuela ; Oosterhoff, Marije ; Frew, Emma ; van Wilder, Philippe ; Postma, Maarten J. ; Evers, Silvia

DOI:

[10.1016/j.hlpt.2020.03.005](https://doi.org/10.1016/j.hlpt.2020.03.005)

License:

Creative Commons: Attribution-NonCommercial-NoDerivs (CC BY-NC-ND)

Document Version

Peer reviewed version

Citation for published version (Harvard):

Schwander, B, Nuijten, M, Hiligsmann, M, Queally, M, Leidl, R, Joore, M, Oosterhoff, M, Frew, E, van Wilder, P, Postma, MJ & Evers, S 2020, 'Identification and expert panel rating of key structural approaches applied in health economic obesity models', *Health Policy and Technology*, vol. 9, no. 3, pp. 314-322.
<https://doi.org/10.1016/j.hlpt.2020.03.005>

[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.

1 Identification and Expert Panel Rating of Key Structural 2 Approaches applied in Health Economic Obesity Models

3 **Authors:** Björn Schwander^{1,2*} (bjoern.schwander@ahead-net.de), Dr. Mark Nuijten³
4 (mark@a2m.nl), Dr. Mickaël Hiligsmann¹ (m.hiligsmann@maastrichtuniversity.nl), Dr.
5 Michelle Queally⁴ (michelle.queally@nuigalway.ie), Prof. Dr. Reiner Leidl^{5,6} ([leidl@helmholtz-
muenchen.de](mailto:leidl@helmholtz-
6 muenchen.de)), Prof. Dr. Manuela Joore⁷ (m.joore@mumc.nl), Marije Oosterhoff⁷
7 (marije.oosterhoff@mumc.nl), Dr. Emma Frew⁸ (e.frew@bham.ac.uk), Prof. Dr. Philippe van
8 Wilder⁹ (philippe.van.wilder@ulb.ac.be), Prof. Dr. Maarten Postma^{10,11,12}
9 (m.j.postma@ruq.nl), Prof. Dr. Silvia Evers^{1,13} (s.evers@maastrichtuniversity.nl)

10 **Acknowledgement:** In addition to the authors Prof. Dr. William Hollingworth¹⁴
11 (william.hollingworth@bristol.ac.uk) was part of the expert panel

12 ***= Corresponding Author**

13 ¹ Department of Health Services Research, CAPHRI - Care and Public Health Research Institute,
14 Maastricht University, Maastricht, The Netherlands

15 ² AHEAD GmbH – Agency for Health Economic Assessment and Dissemination, Lörrach, Germany

16 ³ a2m - Ars Accessus Medica, Amsterdam, The Netherlands

17 ⁴ Department of Economics, National University of Ireland, Galway, Ireland

18 ⁵ Institute of Health Economics and Health Care Management, Helmholtz Zentrum München,

19 ⁶ German Research Center for Environmental Health (GmbH), Neuherberg, Germany

20 ⁷ Department of Clinical Epidemiology and Medical Technology Assessment, Maastricht University
21 Medical Centre+, CAPHRI - Care and Public Health Research Institute, Maastricht, The Netherlands

22 ⁸ Institute of Applied Health Research, University of Birmingham, Birmingham, UK

23 ⁹ Ecole de Santé Publique - Université Libre de Bruxelles, Brussels, Belgium

24 ¹⁰ Unit of PharmacoEpidemiology & PharmacoEconomics, University of Groningen, Groningen, The
25 Netherlands

26 ¹¹ Department of Health Sciences, University of Groningen, University Medical Center Groningen,
27 Groningen, The Netherlands

28 ¹² Department of Economics, Econometrics & Finance, University of Groningen, Faculty of Economics
29 & Business, Groningen, The Netherlands

30 ¹³ Trimbos Institute - Netherlands Institute of Mental Health and Addiction, Utrecht, The Netherlands

31 ¹⁴ School of Social and Community Medicine, University of Bristol, Bristol, UK

32 **Funding:** No funding was provided to assist in the preparation of this research.

33 **Conflict of Interest:** The authors have no other relevant affiliations or financial involvement
34 with any organization or entity with a financial interest in or financial conflict with the
35 subject matter or materials discussed in the manuscript apart from those disclosed.

36 **Abstract (250 of 250 words)**

37 **Objectives:** This study aims to assess the key structural modelling approaches applied in published
38 obesity models, and to provide an expert consensus to improve the methodology and consistency of
39 the application of decision-analytic modelling in obesity research.

40 **Methods:** Using a previously published systematic literature search as basis, ten individual interviews,
41 and a face-to-face expert panel meeting were conducted. Within the expert panel meeting, the
42 interview findings were presented and discussed, rated and where possible consensus statements
43 were obtained. In particular, five topics of interest were assessed: time horizon, model type, obesity-
44 related clinical events simulated, event simulation approaches and external event validation.

45 **Results:** In addition to generic modelling standards, several obesity-specific recommendations were
46 generated: Simulating a lifetime horizon was regarded as optimal (100% agreement); Ideally, both
47 short and long-term results should be presented (100%); Using a risk equation approach for simulating
48 the clinical events was the most preferred approach (60%) followed by applying a body mass index
49 (BMI) related relative risk to a base risk estimate (30%); Continuous BMI approaches were preferred
50 (100%); An individual patient/microsimulation state transition model was regarded as preferred
51 modelling approach (90%); Discrete event simulation (DES) was regarded as the most flexible
52 approach for building an obesity model but it was recognised as complex, and more difficult to build,
53 populate and to disseminate; Performing an external validation was rated as important (100%).

54 **Conclusions:** The obtained insights, discussion and consensus can provide valuable information for
55 developing decision-analytic models to generate high-quality and transparent economic evidence for
56 obesity interventions.

57

58 **Introduction**

59 Obesity, a major public health concern, is a multifactorial disease, caused by both environmental and
60 genetic factors [1], that has reached epidemic proportions globally [2]. The worldwide prevalence of
61 overweight and obesity has doubled since 1980 to an extent that nearly a third of the world's
62 population is now classified as overweight or obese [3]. A common measure to define obesity is the
63 body mass index (BMI), which is obtained by a person's weight in kilograms divided by the square of
64 his height in meters (kg/m^2). According to the WHO definition, a $\text{BMI} \geq 25$ and < 30 in adults is
65 overweight; a $\text{BMI} \geq 30$ in adults is obesity [2].

66 In 2015, high BMI contributed to 4.0 million deaths, which represented 7.1% of the deaths from any
67 cause; it also contributed to 120 million disability-adjusted life-years, which represented 4.9 of
68 disability-adjusted life-years from any cause among adults globally [4] Both overweight and obesity
69 are associated with the incidence of multiple co-morbidities including type II diabetes, cancer and
70 cardiovascular diseases [5].

71 These considerable health impacts of obesity are accompanied with a substantial economic burden,
72 which highlights that there is an urgent need for public health measures in order to save societal
73 resources [6]. Due to this considerable economic impact health economic evaluations are quite
74 commonly applied in the context of obesity prevention and management. Such evaluations allow
75 decision makers to make an informed judgement on the health economic impact of an intervention, by
76 assessing the additional benefits of funding an intervention relative to its additional costs [7]. As shown
77 in systematic reviews [8, 9] decision analytic modelling has commonly been used to evaluate the long-
78 term economic consequences of obesity prevention and therapy measures. In the context of these
79 obesity related decision models the key structural aspects are of fundamental influence as they are
80 impacting all outcomes simulated by the model, including clinical parameters & events, quality of life,
81 direct and indirect costs and hence the whole spectrum of relevant economic consequences [10].

82 Previously, it was shown that there are huge variations in the structural modelling approaches focusing
83 on the prevention and therapy of obesity [8, 9] and up to now no consensus meeting on the structural
84 aspects of obesity models has been performed. This makes it difficult for researchers to select an
85 appropriate approach when designing a model, and subsequently for policy makers and stakeholders
86 to assess the quality of an applied model, intended to inform political or medical decision making.

87 The aim of this study is therefore to assess and measure expert group consensus for key structural
88 modelling approaches of obesity models, and to provide information and recommendations for
89 modellers and decision makers.

90

91 **Methods**

92 On the basis of a previously published systematic literature review [8, 9], the key structural
93 approaches applied in published obesity models were identified.

94 In particular, five inter-related topics of interest were assessed: time horizon, model type, obesity-
95 related clinical events simulated, event simulation approaches and external event validation. These

96 features represent the structural aspects of models listed within the Phillips reporting checklist [11]
97 which are not related on the quality of research reporting (as e.g. statement of the decision problem or
98 statement of scope / perspective etc.). Additionally these features showed a huge variation in
99 published obesity models [8, 9].

100 The findings from the systematic literature review were then used to guide the topic content of the
101 subsequent ten individual interviews. Data from the combined interviews were then presented and
102 discussed at a face-to-face group meeting in order to derive consensus statements with respect to the
103 key structural approaches applied in published obesity models.

104

105 ***Systematic Literature Search***

106 The interviews and the group meeting were informed by a previously published systematic review [8,
107 9] that was performed in the PubMed Database and the NHS Economic Evaluation Database,
108 following the PRISMA guidelines [12]. Three different searches were combined: one for health
109 economic evaluations, one for decision models and one for obesity. Eligible studies were original
110 research articles on decision models for full health economic assessment in the context of obesity; the
111 definitions from Drummond et al. [13] (health economic assessments), from the International Society
112 for Pharmacoeconomics and Outcomes Research (ISPOR) Task Force [14] (decision models), and
113 from the WHO criteria [2] (obesity) were applied in order to define eligible studies. In total 4,293
114 studies were identified via the database searches, and were reviewed. From these 142 articles were
115 selected for full-text review; of which 87 papers met the inclusion criteria. Of those, 72 models
116 simulated obesity associated events. The rationale for this selection is on one hand to investigate the
117 selected event simulation approaches. On the other hand the rationale is to enable performing and
118 investigating external validations of the event projections made by these models, which increases the
119 credibility of the modelling approaches for researchers, physicians and decision makers.

120 For more details on the literature search, the eligibility criteria and the literature selection please refer
121 to the published systematic review [8, 9].

122

123 ***Individual interviews***

124 Several health economic experts, with in-depth experience in decision analytic modelling and/or
125 economics of obesity (using a convenience sampling), were requested to participate in an Expert

126 panel meeting during the European Health Economic Association (EuHEA) conference 2018 in
127 Maastricht, and ten (of twenty-two contacted) agreed to participate the meeting and to perform a 60-
128 minute individual preparation interview beforehand. Within this interview the outcomes of the
129 previously published systematic review, related to the key structural aspects (time horizon, model type,
130 obesity-related clinical events simulated, event simulation approaches and external event validation)
131 were presented via a web-based platform, and related to each of the key structural aspects of a model
132 specific questions were asked.

133 With respect to the choice of a specific event simulation approach, different definitions were first
134 obtained from the systematic review [9] and are presented, together with the interview questions, in
135 box 1.

136

137 *Please add box 1 here*

138

139 The individual interview data were then analysed quantitatively in MS Excel and summarized in a MS
140 PPT presentation in order to serve as basis for the discussions at the expert panel meeting.

141

142 ***Expert Panel Meeting***

143 The face-to-face expert panel meeting was performed as satellite event of the EuHEA conference in
144 Maastricht, on July 13th 2018. Within this meeting, the interview results relating to each question were
145 presented and discussed, with the aim of reaching a group consensus or to capture the variance in
146 opinion for each item. Within this meeting the key structural aspects, were discussed in detail with a
147 specific focus on obesity-specific criteria. After the meeting the results were summarized and sent to
148 the expert panel members for further comment and approval.

149 The results from this expert panel meeting are presented below, together with the results of the
150 individual interviews and the key results from the systematic literature review.

151

152 **Results**

153 ***Time Horizon***

154 Table 1 presents the outcomes linked to the choice of time horizon for all published models identified
155 in the review, and for the expert group opinion.

156

157 *Please add table 1 here*

158

159 In the expert panel meeting, it was agreed that a lifetime horizon is optimal for a health economic
160 obesity model (100% agreement)) and it was further agreed that both short- and long-term results
161 should be presented (100% agreement). Short-term / trial period simulations may indeed also be
162 interesting for practitioners / physicians, and are less susceptible to assumptions such as the
163 sustainability of the intervention effect size and the natural course / development of BMI over time,
164 including potential weight-regain post intervention.

165

166 ***Obesity Associated Events***

167 Table 2 illustrates the findings from the literature review with respect to obesity-associated events
168 (based on the 72 studies that have simulated obesity-associated events) alongside the findings from
169 the expert interviews. Most of the published models simulated coronary heart disease (CHD) ($\approx 83\%$;
170 60 of 72), type 2 diabetes (T2D) ($\approx 74\%$), and stroke ($\approx 67\%$). A minority of the models simulated
171 cancer ($\approx 35\%$), osteoarthritis ($\approx 24\%$), hyperlipidaemia ($\approx 11\%$), hypertension ($\approx 11\%$), and peripheral
172 arterial disease ($\approx 10\%$).

173

174 *Please add table 2 here*

175

176 From the expert interviews, with regard to the question on the minimum acceptable events to be
177 included in a health economic obesity model (presented in table 2), in 50% of cases only CHD, T2D
178 and stroke were named as “minimum acceptable events” in 20% of cases accompanied by cancer and
179 in 10% accompanied by hypertension; whereas in two cases no definite answer was given due to the
180 rationale that “in general those events with strongest association / causal relationship to obesity should
181 be included”. Related to the question on the events to be included in a health economic obesity model
182 in the optimal world (presented in Figure 1) the picture was more diverse.

183

184 *Please add figure 1 here*

185

186 In 40% of cases it was stated that all events with a clear association with obesity should be included.
187 One expert stated that this clear association should be combined with the severity of event
188 consequences. In 50% of cases, CHD, T2D and stroke were named (alone or in combination with
189 other diseases), whereas by one expert no definite answer was given as it was claimed that it depends
190 on the goal of the model and on the available evidence.

191 During the expert panel, several discussions around these obesity associated events took place
192 (please refer to discussion part), but it was not possible to achieve consensus on the whole. However,
193 finally there was general agreement that those events with a strong statistical association to obesity
194 combined with a clear clinical causal relationship to obesity should be included in the optimal case.

195

196 **Model Type**

197 Table 3 presents the results concerning the appropriate model type.

198

199 *Please add table 3 here*

200

201 In the expert interviews, in 90% of cases a state transition model was named as the preferable
202 approach, and, within these responses - 60% suggested a state transition model alone, and 30% also
203 recommended a DES as an alternative model type to consider. Only one expert (10%) recommended
204 DES alone.

205 On the question “why a specific model type was preferred?” the following rationales were provided by
206 the experts:

- 207 • *“STM is adequate to simulate the three major health impacts (T2D, CHD and stroke);*
- 208 • *STM is most practicable for event based simulation;*
- 209 • *STM is the most familiar approach (for health economists and stakeholders);*
- 210 • *STM is the most familiar approach - and individual patient simulation enables; building in*
211 *specific memory;*
- 212 • *An individual patient simulation STM is preferred as it is possible to include a kind of memory”.*

213 In three cases both the DES and the STM were preferred by the experts, for the following reasons:

- 214 • *“Memory is an important factor (as time with obesity / related morbidity impacts event risk) -*
215 *therefore a DES would be preferred or a STM on a patient level with included memory states;*

- 216 • *Due to competing risks a DES / STM using a microsimulation approach will be preferred (for*
217 *DES not all data might be available);*
- 218 • *DES might be scientifically the best approach but difficult to build, inform and to explain. STM*
219 *might be the most accepted approach”.*

220 For one participant the DES alone was preferred as

- 221 • *“DES allows considering timing of events which is important due to the inter-event*
222 *dependencies”.*

223 Within the expert panel, a consensus was reached in the form of the following two statements:

- 224 • *An individual patient / microsimulation STM is regarded as preferred approach for an obesity*
225 *model;*
- 226 • *DES is regarded as the most flexible approach however DES is complex, difficult to build, to*
227 *inform and to explain (to stakeholders).*

228

229 ***Event Simulation Approach***

230 Within the expert interviews the experts were asked to rank a list of potential modelling approaches
231 identified from the systematic review. The results are presented in table 4 and in figure 2,
232 respectively. The risk equation approach was the most preferred approach (60% rated this as number
233 one, followed by BMI-related RR (30% rated this as number one) and one expert felt it difficult to rank
234 the approaches.

235

236 *Please add table 4 here*

237

238 *Please add figure 2 here*

239 The reasons for the number one rating for the Equation / Change in Risk Factors were:

- 240 • *“Method is quite robust, widely validated and widely used;*
- 241 • *Quite valid (accepted) approach and most commonly used;*
- 242 • *Not everything might be explainable by change in BMI and therefore it may be important to*
243 *consider further risk factors;*
- 244 • *Risk equation approach describes the whole nature of a chronic disease;*
- 245 • *Risk equation approach takes into account inter-event dependencies;*

- 246 • *Risk equation approach is widely applied and health economists are most familiar with this;*
247 • *Familiar approach, well know, risk equations are also used in clinical guidelines; for the others*
248 *it is the key question how strong the association between BMI and risk is”.*

249 The reasons for the number one rating for the Incidence / BMI related RR were:

- 250 • *“Most valuable / simple to set up events driven models for obesity;*
251 • *BMI related RR is preferred as always small changes are taken into account;*
252 • *Continuous BMI approaches are preferred against categorical approaches (there was 100%*
253 *agreement on this statement in the expert panel)”.*

254

255 Furthermore, in the interviews, the experts were asked whether they would suggest using different
256 approaches for different events if considering CHD, T2D, and stroke. With regard to this question, 90%
257 answered with “no”; whereas 40% mentioned that not necessarily different approaches need to be
258 applied and 50% answered that consistent approaches (if applicable) are preferred. One expert found
259 it difficult to rate this topic and gave no answer.

260

261 ***External Validation***

262 External validation was defined as comparing the model's results with actual event data [15]. External
263 validation involves simulating events that have occurred, such as those in clinical trials or
264 epidemiologic studies, and examining how well the model results compare.

265 According to the systematic review, only ten published model-based health economic assessments in
266 obesity included an external event validation (14%; 10 of 72).

267 Within the individual interviews the experts were asked how important they rate an external validation
268 with possible answers being: “essential”, “very important”, “important”, “less important”, “not important”
269 or “other” (please specify). All experts (100%) rated the external validation as “important”; 60% “very
270 important” and 20% as “essential”. These findings were confirmed during the expert panel.

271

272 ***Summary of Key Recommendations***

273 A summary of key recommendations generated as a result of the expert interviews combined with the
274 expert panel meeting are presented in table 5.

275

276 *Please add table 5 here*

277

278 **Discussion**

279 Focusing on the key structural aspects outlined in the Philips checklist [11], this paper presents the
280 main findings relevant to obesity models that have been identified (systematic literature search), rated
281 (expert interviews) and discussed (expert panel). The expert panel meeting resulted in specific
282 modelling recommendations that go beyond the findings from the systematic literature research, which
283 is also representing the novelty of this research. The main findings by key structural aspect are
284 discussed in detail below; each topic starts with a summary of outcomes of the expert panel meeting
285 and these outcomes are then discussed and set into perspective by reflecting the complex
286 circumstances and considerations related to each aspect. The latter discussion points are mainly
287 driven by statements obtained during the expert panel meeting, which were accompanied and
288 completed on the basis of related literature.

289

290 ***Time Horizon***

291 With regard to the time horizon of a health economic obesity model, it was possible to obtain clear
292 expert recommendations. However, there were some interesting viewpoints expressed during the
293 expert panel mostly around the question of whether or not a short term (e.g. trial period) simulation
294 should be performed and presented. One key consideration in this context was that practitioners,
295 physicians and stakeholders might be (additionally) interested in short term results and it is
296 recommended that health economists also take into account the information needs of the health care
297 personnel involved and also the requests / preferences of policy makers and other stakeholders. From
298 a scientific point of view the key reasons for presenting short term / trial period outcomes (in addition
299 to lifetime) were to present the impact of lifetime extrapolations as well as the practical need to
300 determine whether the model adequately replicates the underlying study/trial results (internal
301 validation). The key issues of extrapolation named in the context of obesity were the sustainability of
302 the effect size (e.g. weight or BMI reduction and the related regain over time) and the natural
303 course/development of weight / BMI over time, which is often based on a limited time-horizon, which
304 again requires extrapolation to lifetime. These key issues of extrapolation were the key drivers for
305 recommending an additional presentation of short term / trial period results.

306 **Obesity Associated Events**

307 The discussions around obesity-associated events to be modelled reflected some divergent views but
308 there was general alignment among the experts that those events with a strong association to obesity
309 combined with a clear causal relationship to obesity should be included in the optimal case. In contrast
310 to the causal relationship of a specific event the strength of association could be more easily
311 assessed, as the odds ratio or relative risk based on the best case could be extracted from
312 prospective cohort studies. In a systematic review and meta-analysis of Guh et al. 2009 [5] the
313 relative risk of various obesity associated events was presented and results by prospective cohort
314 study and pooled results were provided, by gender and weight status (overweight / obese). According
315 to the pooled results for obesity the strongest RR based associations in females (defined as $RR \geq 2$ in
316 subjects with a $BMI \geq 30$) were obtained for T2D ($RR=12.41$), CHD ($RR=3.10$), Gallbladder Disease
317 ($RR=3.08$), Endometrial Cancer ($RR=2.86$), Kidney Cancer ($RR=2.64$), Hypertension ($RR=2.42$),
318 osteoarthritis ($RR=2.19$) and congestive heart failure ($RR=2.06$) [5]. For males the strongest RR based
319 associations (defined as $RR \geq 2$ in subjects with a $BMI \geq 30$) were obtained for T2D ($RR=6.74$),
320 osteoarthritis ($RR=4.20$), pancreatic cancer ($RR=2.29$) and asthma ($RR= 2.19$); the association to
321 CHD in males ($RR=1.75$) was not that pronounced as in females ($RR=3.10$) [5]. Furthermore the
322 association of obesity and stroke was not that pronounced with a RR of 1.50 in females and a RR of
323 1.68 in males [5]. Hence looking at the results of the systematic review (T2D, CHD and stroke are the
324 most frequently included events within health economic obesity models) it is clear that not only the
325 strength of association is important but also the severity and consequences of the specific events
326 need to be considered, which was also discussed and determined as a selection criteria during the
327 expert panel meeting, and might explain the brought inclusion of CHD and stroke into the health
328 economic obesity models, as both events are potentially leading to mortality or disability. Furthermore
329 from a health economic perspective the absolute incidence of events plays a role, as a strong obesity-
330 association that is observed only in a very small number of patients, might have less impact on the
331 cost-effectiveness than an event with a weak obesity-association that is observed in many patients.

332 The answer on the strength of statistical association, the severity and the absolute incidence of events
333 are much easier to be answered than the question on the causal clinical relationship to obesity. The
334 passage from obesity to T2D is caused by a progressive defect in insulin secretion coupled with a
335 progressive rise in insulin resistance. Both insulin resistance and defective insulin secretion appear

336 very prematurely in patients with obesity, and both worsen similarly towards diabetes [16], therefore
337 the causal relationship is well understood. Also, there is good evidence on the causal relationship
338 between obesity and CHD, and obesity and stroke and insulin resistance has been identified as the
339 primary mechanism driving the progression of cardio-metabolic diseases (such as CHD and stroke)
340 [17]. For different types of cancer the causal relationship is more challenging to capture and it remains
341 unclear how obesity impacts the aetiology of cancer, which itself is not fully understood [18]. Hence,
342 many researchers might have not included cancer as an obesity associated event within the model. If
343 including only those events, for which there is clear evidence of a causal relationship, T2D, CHD and
344 stroke would be an adequate minimum selection to be simulated within a health economic model. In
345 this context it is recommended that the inclusion of events for which the causal relationship to obesity
346 is not yet fully understood is investigated within scenario analyses.

347

348 **Model Type**

349 The model types recommended for a health economic obesity model were either an individual patient /
350 microsimulation STM or alternatively a DES. DES is clearly understood as the most flexible approach
351 for building an obesity model, but it was also recognised as complex, as more difficult to build,
352 populate and to disseminate (to stakeholders). Many shortcomings of (cohort) state transition models
353 can be compensated by an individual patient / microsimulation approaches which enables patient
354 history to be tracked using tunnel states and therefore overcome the Markovian assumption; this is
355 important for obesity as time with obesity and/or obesity associated morbidities impacts the event risk.
356 However, there is still some functionality of DES models that cannot be reproduced by a STM [19].
357 The DES can simulate interactions amongst individuals or between individuals and the environment
358 [20, 21], which might be interesting in obesity prevention models in which the positive effect of an
359 intervention could have a positive effect on the whole community (e.g. on a whole school class or the
360 whole school setting). Furthermore DES is well suited to modelling situations where patients are
361 subject to multiple or competing risks [21, 22]. A DES manages the competing and the sequencing of
362 events by generating a future events list, then, for example, selecting the next closest time-to-event to
363 ascertain which event occurs next in the process. This is relevant for obesity as there are several
364 obesity associated events to be simulated. In a STM a transition probability is derived for each
365 mutually exclusive competing health state and these competing health states must be exhaustive, and

366 it requires many health states to achieve a level of detail comparable to DES. In a DES it is also easier
367 to manage multiple events at the same time and to include and exclude events [23]. In the STM the
368 patient is in one of a variety of mutually exclusive health states at any one time, which need to be
369 clearly defined in the model structure, hence including / excluding events is a complex task.
370 Furthermore, DES models can capture a greater level of detail than STM allowing the model to
371 capture more detail regarding uncertainty in the system and including time to event information [21,
372 22]; this is important for obesity as multifactorial conditions and complex interventions (e.g. in the
373 context of prevention) need to be simulated.

374 Besides all these advantages it needs to be considered that there are also several disadvantages,
375 which prevent a broad application of DES in the fields of health economics [19]. DES models are
376 generally more complex, require more data (that is often not available), and take more time to develop
377 and run than STM; furthermore this could lead to a DES-induced over-specification [24] where models
378 may become more complex than necessary, which again leads to increased data needs for DES
379 models compared to STM [24].

380 These issues prevent a broad application of DES in health economics of obesity. The STM is rated as
381 a pragmatic, widely applied, practical, familiar and widely accepted approach by the expert panel.
382 Especially the communication and dissemination of (complex) DES models to stakeholders and policy
383 makers is seen as a key hurdle for a broad application, as usually the model approach needs to be
384 understandable to achieve research impact.

385

386 ***Clinical Event Simulation***

387 The obtained event simulation approaches are quite diverse but it was possible to identify two
388 preferred approaches by the expert panel namely the risk equation approach (most preferred
389 approach - 60% rated this as number one, and the BMI related RR (30% rated this as number one).

390 Many reasons were provided by the experts why the risk equation approach is preferred. The most
391 prominent ones were that the risk equation approach describes the whole nature of a chronic disease
392 and takes into account inter-event dependencies whereas within the BMI based approach the question
393 remains whether everything can be explained only by the BMI and how strong the BMI association of a
394 specific disease really is. A further point that was highlighted in the expert discussions was that the
395 modellers' decision on the event simulation approach is often driven by data availability. Whereas for

396 the BMI based approach only data on the BMI development (over time) is required, the risk equation
397 approach requires data on all risk factors included in the equation, and is therefore far more data
398 demanding. In the case that data on the risk factors is not available the BMI approach could be the
399 most pragmatic way to estimate the health economic impact of an intervention, although the named
400 limitations need to be considered and extra sensitivity analyses and scenario analysis may be
401 required. Furthermore the experts agreed on the procedure that (if possible) comparable event
402 simulation approaches should be applied for the different events, mainly to have comparable strengths
403 and limitations for the simulation of the different events included in the obesity model.

404

405 ***External Validation***

406 The systematic review identified only ten models (of 72 that simulated events) that performed an
407 external validation [9]. As this procedure is a key part of testing the validity of the modelling results
408 with regard to the predictiveness of the event simulation approach, this was in general regarded as a
409 limitation of published obesity models. All the experts rated the external validation as (at least)
410 important for a health economic obesity model and that this should be performed as standard together
411 with the internal validation that is usually performed as part of the internal model testing.

412

413 ***General Issues of Obesity Models***

414 Besides the key structural aspects that were investigated and discussed there are several other
415 aspects that make it a challenge to model health economic assessments in obesity. As already
416 mentioned one key difficulty is that the chronic events associated with obesity require a lifetime
417 horizon and therefore several assumptions related to the sustainability of the effect size and the
418 natural course of weight / BMI. It is recommended that these two factors require clear and transparent
419 handling and need to be investigated in a sensitivity analyses.

420 One other aspect that makes obesity models so diverse is that an intervention might focus either on
421 the therapy or on the prevention of obesity. Whereas prevention measures usually start in younger
422 age groups (e.g. in the school setting), the therapy of obesity could either target young or older age
423 populations. Modelling prevention measures are usually more complex than modelling therapy, as the
424 prevention effect might have a positive influence on the whole community setting, and would hence

425 require simulating interactions amongst individuals or between individuals and the environment,
426 whereas therapy is usually targeted to the patients receiving a specific intervention.

427 Besides the diversity in the setting and intervention there are quite some challenges related to the
428 understanding of the aetiology of obesity and of obesity associated diseases including so called
429 obesity-paradoxes [25]. Whereas obesity implies increased risk for chronic diseases, it is in fact
430 associated with decreased mortality risk compared with normal weight [25]. Another paradox concerns
431 the observation that when fitness is taken into account, the mortality risk associated with obesity is
432 offset [25]. Furthermore there is a paradox describing the presence of a sizeable subset of individuals
433 with obesity who are otherwise healthy [25]. Even when some obese persons are healthy and for late
434 phase of disease, obesity may be protective, it still is considered an important risk factor in the
435 development of chronic disease. This has been recently stressed in a review on cardiovascular
436 diseases [26]. Modelling may thus have to distinguish several subgroups, depending on time and
437 diseases analysed.

438

439 ***Limitations and Implications***

440 As discussed above, challenges around the economic modelling of obesity are not purely structural,
441 and hence one limitation of this study is the focus only on key structural aspects. However, especially
442 as there are many challenges, it is important to offer recommendations on the handling of some key
443 structural aspects when simulating obesity. The rationale for this is that the basic structure of the
444 model is integral, and each decision that is made in the key structural development is carried forward
445 to each calculation step of the model. Therefore the provided consensus on those fundamental
446 structural issues could minimize the challenges modellers, stakeholders and decision makes face,
447 while developing, interpreting and rating model-based health economic assessments in obesity.

448 For the expert panel, we focused on experts that were attending the EuHEA meeting in Maastricht
449 (2018), as a result of this selection criterion we had only European experts participating. Hence one
450 limitation of this approach was that researches from non-European countries were not able to
451 contribute to this research. Considering that, according to the previously published systematic review,
452 47% of decision models focused on a European setting, 27% on US setting and 20% on an Australian
453 setting, it would have been interesting to consider additionally the expert opinion of non-European
454 experts. Additionally the limited number of experts (n=10 experts) has to be rated as limitation of our

455 research, therefore it could be interesting to validate our findings on the basis of a larger sample of
456 experts. Accordingly the presented work is not to be seen as a strict guideline for obesity modelling but
457 as a set of information and recommendations that are regarded to be useful for other researchers and
458 decision makers in this field.

459 Further, in the expert interviews and in the expert panel we only used basic quantitative methods in
460 order to obtain an expert rating and an expert consensus, as the style of questions were not designed
461 to involve more advanced quantitative methods (e.g. discrete choice experiments) or qualitative
462 techniques (such as the Delphi method). Furthermore the set focus on health economists is a
463 limitation related to the composition of the panel. The rationale for selecting health economists was
464 that modelling is primarily driven by this discipline, but as a consequence it was not possible to get a
465 clear expert rating on purely clinical aspects, such as the obesity associated event selection. In case
466 of specialized epidemiologists and / or clinicians the discussion might have moved more into the
467 direction of which events are nowadays considered as clearly obesity associated, a fact that we have
468 tried to resolve by discussing the latest related literature.

469 Although we have observed consensus on many structural issues, there is no structural approach that
470 covers all needs, and hence related to the decision problem, research question, and according to the
471 data and resource availability there are different structural approaches that were rated as suitable for
472 building a health economic obesity model. Furthermore, depending on the purpose of a health
473 economic evaluation in obesity, researches might take different approaches than those recommended
474 in our paper, if they have a good rationale for doing so.

475 One key question that remains in this context is, how the application of different approaches to the
476 same decision problem, research question and population might influence the results of the clinical
477 event prediction and subsequently of the whole health economic evaluation – which is seen as a
478 valuable field of future research.

479

480 **Conclusions**

481 While the working group acknowledges the challenges in achieving consensus, several
482 recommendations for the key structural approaches for a health economic obesity model were
483 developed. The obtained insights, discussion content and consensus can provide valuable information

484 for all decision makers, health economists and modellers for developing decision-analytic models to
485 generate high-quality and transparent economic evidence for obesity interventions.

486

487 **List of Abbreviations**

Abbreviation	Meaning
BMI	Body mass index
CHD	Coronary heart disease
DES	Discrete Event Simulation
EuHEA	European Health Economic Association
HEA	Health Economic Assessment
ISPOR	International Society for Pharmacoeconomics and Outcomes Research
PRISMA	Preferred reporting items for systematic reviews and meta-analyses
RR	Relative risk
STM	State transition model
T2D	Type 2 diabetes
UKPDS	United Kingdom Prospective Diabetes Study
WHO	World Health Organization

488

489

490 **References**

- 491 1. Huang T, Hu FB: **Gene-environment interactions and obesity: recent developments and**
492 **future directions.** *BMC medical genomics* 2015, **8 Suppl 1**:S2.
- 493 2. World Health Organization. **Fact Sheet on Obesity.** 2003.
494 [https://www.who.int/dietphysicalactivity/media/en/gsf_obesity.pdf] (Accessed on
495 09.02.2019).
- 496 3. Chooi YC, Ding C, Magkos F: **The epidemiology of obesity.** *Metabolism: clinical and*
497 *experimental* 2019, **92**:6-10.
- 498 4. Afshin A, Forouzanfar MH, Reitsma MB *et al.* **Health Effects of Overweight and Obesity in**
499 **195 Countries over 25 Years.** *The New England journal of medicine* 2017, **377**(1):13-27.
- 500 5. Guh DP, Zhang W, Bansback N *et al.* **The incidence of co-morbidities related to obesity**
501 **and overweight: a systematic review and meta-analysis.** *BMC public health* 2009, **9**:88.
- 502 6. Tremmel M, Gerdtham UG, Nilsson PM, Saha S: **Economic Burden of Obesity: A**
503 **Systematic Literature Review.** *International journal of environmental research and public*
504 *health* 2017, **14**(4).
- 505 7. Ananthapavan J, Sacks G, Moodie M, Carter R: **Economics of obesity--learning from the**
506 **past to contribute to a better future.** *International journal of environmental research and*
507 *public health* 2014, **11**(4):4007-4025.
- 508 8. Schwander B, Hiligsmann M, Nuijten M, Evers S: **Systematic review and overview of health**
509 **economic evaluation models in obesity prevention and therapy.** *Expert review of*
510 *pharmacoeconomics & outcomes research* 2016, **16**(5):561-570.
- 511 9. Schwander B, Nuijten M, Hiligsmann M, Evers S: **Event simulation and external validation**
512 **applied in published health economic models for obesity: a systematic review.** *Expert*
513 *review of pharmacoeconomics & outcomes research* 2018, **18**(5):529-541.
- 514 10. Afzali HH, Karnon J, Merlin T: **Improving the accuracy and comparability of model-based**
515 **economic evaluations of health technologies for reimbursement decisions: a**
516 **methodological framework for the development of reference models.** *Medical decision*
517 *making : an international journal of the Society for Medical Decision Making* 2013, **33**(3):325-
518 332.
- 519 11. Philips Z, Bojke L, Sculpher M, Claxton K, Golder S: **Good practice guidelines for decision-**
520 **analytic modelling in health technology assessment: a review and consolidation of**
521 **quality assessment.** *PharmacoEconomics* 2006, **24**(4):355-371.
- 522 12. Moher D, Liberati A, Tetzlaff J, Altman DG: **Preferred reporting items for systematic**
523 **reviews and meta-analyses: the PRISMA statement.** *Journal of clinical epidemiology* 2009,
524 **62**(10):1006-1012.
- 525 13. Drummond MF, Sculpher MJ, Claxton K, Stoddart GL, Torrance GW: **Methods for the**
526 **Economic Evaluation of Health Care Programmes.** Oxford: Oxford: Oxford University
527 Press; 2015.
- 528 14. Weinstein MC, O'Brien B, Hornberger J *et al.* **Principles of good practice for decision**
529 **analytic modeling in health-care evaluation: report of the ISPOR Task Force on Good**
530 **Research Practices--Modeling Studies.** *Value in health : the journal of the International*
531 *Society for Pharmacoeconomics and Outcomes Research* 2003, **6**(1):9-17.
- 532 15. Eddy DM, Hollingworth W, Caro JJ *et al.* **Model transparency and validation: a report of**
533 **the ISPOR-SMDM Modeling Good Research Practices Task Force--7.** *Value in health : the*

- 534 *journal of the International Society for Pharmacoeconomics and Outcomes Research* 2012,
535 **15(6):843-850.**
- 536 16. Golay A, Ybarra J: **Link between obesity and type 2 diabetes.** *Best practice & research*
537 *Clinical endocrinology & metabolism* 2005, **19(4):649-663.**
- 538 17. Aslibekyan S, Garvey WT: **Obesity: Obesity and cardiometabolic disease - more than**
539 **meets the eye.** *Nature reviews Endocrinology* 2017, **13(10):566-568.**
- 540 18. Stone TW, McPherson M, Gail Darlington L: **Obesity and Cancer: Existing and New**
541 **Hypotheses for a Causal Connection.** *EBioMedicine* 2018, **30:14-28.**
- 542 19. Standfield L, Comans T, Scuffham P: **Markov modeling and discrete event simulation in**
543 **health care: a systematic comparison.** *International journal of technology assessment in*
544 *health care* 2014, **30(2):165-172.**
- 545 20. Brennan A, Chick SE, Davies R: **A taxonomy of model structures for economic evaluation**
546 **of health technologies.** *Health economics* 2006, **15(12):1295-1310.**
- 547 21. Karnon J, Stahl J, Brennan A *et al*: **Modeling using discrete event simulation: a report of**
548 **the ISPOR-SMDM Modeling Good Research Practices Task Force--4.** *Value in health : the*
549 *journal of the International Society for Pharmacoeconomics and Outcomes Research* 2012,
550 **15(6):821-827.**
- 551 22. Caro JJ, Briggs AH, Siebert U, Kuntz KM: **Modeling good research practices--overview: a**
552 **report of the ISPOR-SMDM Modeling Good Research Practices Task Force--1.** *Value in*
553 *health : the journal of the International Society for Pharmacoeconomics and Outcomes*
554 *Research* 2012, **15(6):796-803.**
- 555 23. Caro JJ, Moller J, Getsios D: **Discrete event simulation: the preferred technique for health**
556 **economic evaluations?** *Value in health : the journal of the International Society for*
557 *Pharmacoeconomics and Outcomes Research* 2010, **13(8):1056-1060.**
- 558 24. Karnon J, Brown J: **Selecting a decision model for economic evaluation: a case study**
559 **and review.** *Health care management science* 1998, **1(2):133-140.**
- 560 25. McAuley PA, Blair SN: **Obesity paradoxes.** *Journal of sports sciences* 2011, **29(8):773-782.**
- 561 26. Elagizi A, Kachur S, Lavie CJ *et al*: **An Overview and Update on Obesity and the Obesity**
562 **Paradox in Cardiovascular Diseases.** *Progress in cardiovascular diseases* 2018, **61(2):142-**
563 **150.**
- 564

565 **Box 1: Interview Questions and Definitions of Event Simulation Approaches**

566 **Interview Questions:**

- 567 • Which time horizon would you rate as the minimum acceptable for a health economic
568 obesity model?
- 569 • Which time horizon would you rate as optimal for a health economic obesity model?
- 570 • Which (obesity associated) events would you rate as the minimum acceptable to be
571 included into a health economic obesity model?
- 572 • Which (obesity associated) events would you rate as optimal to be included into a health
573 economic obesity model?
- 574 • Which model type would you prefer for a health economic obesity model?
- 575 • Why would you prefer this model type?
- 576 • Which event simulation approach would you prefer for a health economic obesity model?
577 Please rank the top 3 approaches that you would prefer (1 = most preferred one to 3 =
578 least preferred but still preferred one)
- 579 • Why would you prefer the top rated (#1) event simulation approach?
- 580 • Would you suggest to use different approaches for different events (consider coronary
581 heart disease, type 2 diabetes, stroke)? If yes – why?
- 582 • How important do you rate an external validation for a health economic obesity model?

580 **Definitions of Event Simulation Approaches**

- 581 • Risk Equation / Change in Risk Factors: E.g. Framingham / UKPDS equations – the
582 base risk is calculated as an equation of risk factors and the intervention effect is
583 simulated by the change of risk factors
- 584 • Disease Incidence Estimate / BMI related relative risk (RR): Any kind of incidence
585 estimate (e.g. age-specific; gender-specific incidence etc.) is used as base risk and the
586 intervention effect is simulated by applying a BMI related relative risk to the base risk
- 587 • BMI Function / Change in BMI: Base risk is calculated as function of the BMI which is
588 directly influenced by the intervention effect on the BMI
- 589 • Disease Incidence Estimate / Obesity related RR: Any kind of incidence estimate (e.g.
590 age-specific; gender-specific incidence etc.) is used as base risk and the intervention
591 effect is simulated by applying an obesity status related relative risk (e.g. BMI <30 non-
592 obese ; BMI \geq 30 obese) to the base risk
- 593 • BMI Group Function / Change in BMI Group: Base risk is calculated as function of
594 specific BMI groups (e.g. < 25 normal weight; 25-30 overweight; 30-35 moderate obese;
595 \geq 35 severe obese etc.) which is directly influenced by the intervention effect on the BMI
group
- 596 • Disease Incidence Estimate / BMI Group related RR: Any kind of incidence estimate
(e.g. age-specific; gender-specific etc.) is used as base risk and the intervention effect is
simulated is simulated by applying a BMI group related relative risk to the base risk

596 **Tables**597 **Table 1: Time Horizon – Systematic Literature Search and Expert Interview Outcomes**

Time Horizon	Literature Review (n=87 models)	Expert Interviews (n=10 experts)	
		Minimum	Optimal
< 20 years	23%	20%	10%*
≥ 20 and < lifetime	14%	20%	10%*
Lifetime	63%	60%	100%*

598 * 2 experts provided 2 different answers: ≥ 20 years in adults / lifetime in younger subjects; ≥
599 10 years / lifetime optimal

600

601 **Table 2: Obesity Associated Events – Systematic Literature Search and Expert**
602 **Interview Outcomes**

Obesity Associated Events	Literature Review* Outcomes (n=72 models)	Expert Interviews Outcomes (n=10 experts) (Minimum acceptable events)*		
		ChD, T2D and, Stroke	ChD, T2D, Stroke and Cancer	ChD, T2D, Stroke, Cancer and HT
Coronary heart disease (ChD)	83%	50%*	20%*	10%*
Type 2 Diabetes (T2D)	74%			
Stroke	67%			
Cancer	35%			
Hypertension (HT)	11%			
Osteoarthritis	27%			
Hyperlipidaemia	11%			
Peripheral arterial disease	10%			

603 *no definite answer was provided by 2 experts (n=20%) - in general those events with
604 strongest association / causal relationship to obesity should be included

605

606 **Table 3: Model Type – Systematic Literature Search and Expert Interview Outcomes**

Model Type	Literature Review (n=87 models)	Expert Interviews (n=10 experts)
State Transition Model (STM)	85%	60%
Disease Event Simulation (DES)	2%	10%
Decision Tree Model	13%	--
STM or DES (expert rating)	--	30%

607 * 3 experts rated both STM and DES as suitable - depending on the data availability (for the
608 DES model)

609

610

611 **Table 4: Event Simulation Approach – Systematic Literature Search and Expert**
 612 **Interview Outcomes**

Event Simulation Approach	Literature Review (n=72 models)	Expert Interviews (n=10 experts) – Ranking (#1, #2, #3)
Risk Equation / Change in Risk Factors	32%	#1 (60%); #2 (10%); #3 (20%)
Disease Incidence Estimate / BMI related relative risk (RR)	21%	#1 (30%); #2 (40%); #3 (0%)
BMI Function / Change in BMI	12%	#1 (0%); #2 (20%); #3 (20%)
Disease Incidence Estimate / Obesity related RR	12%	
BMI Group Function / Change in BMI Group	9%	
Disease Incidence Estimate / BMI Group related RR	7%	
Others / Others	7%	

613 * 3 experts rated both STM and DES as suitable - depending on the data availability (for the
 614 DES model)

615

616 **Table 5: Overview of key expert recommendations by key structural aspect**

Key Structural Aspect	Expert panel recommendations
Time Horizon	Simulating a lifetime horizon was regarded as optimal for an obesity model (100% agreement) Ideally, both short and long-term results should be presented (100% agreement)
Obesity Associated Events	No consensus was possible on which clinical events to be included in a health economic obesity model There was general alignment that those events with a strong association to obesity combined with a clear causal relationship to obesity should be included in the optimal case
Model Type	An individual patient/microsimulation state transition model was regarded as preferred modelling approach (90% agreement) Discrete event simulation (DES) was regarded as the most flexible approach for building an obesity model but DES was recognised as complex, as more difficult to build, populate and to disseminate (to stakeholders)
Event Simulation Approach	Using a risk equation approach for simulating the clinical events was the most preferred approach (60%) followed by applying a body mass index (BMI) related relative risk to a base risk estimate (30%) Continuous BMI approaches were preferred (relative to categorical ones) (100% agreement)
External Validation	100% of experts rated the external validation at least important

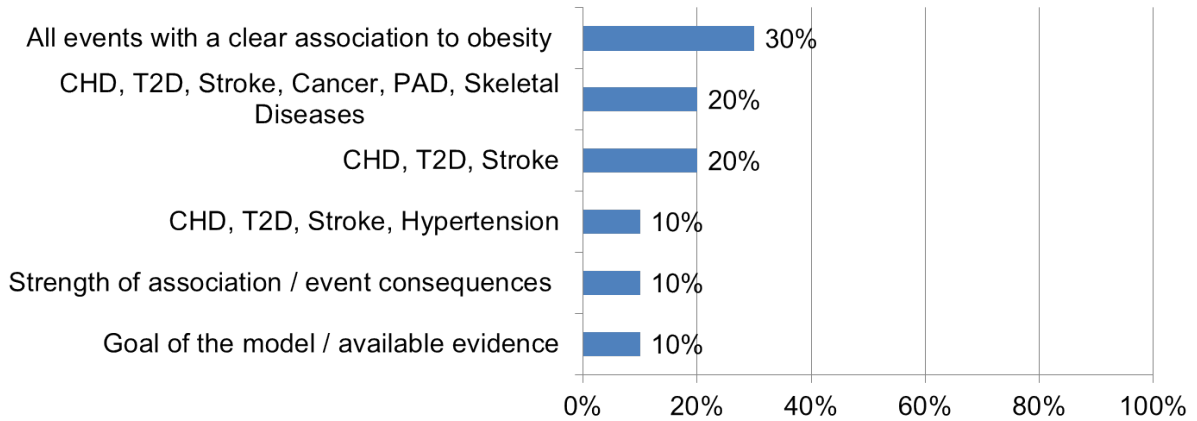
617

618 **Figures**

619

620

621 **Figure 1: Which (obesity associated) events would you rate as optimal to be included**
 622 **into a health economic obesity model?**

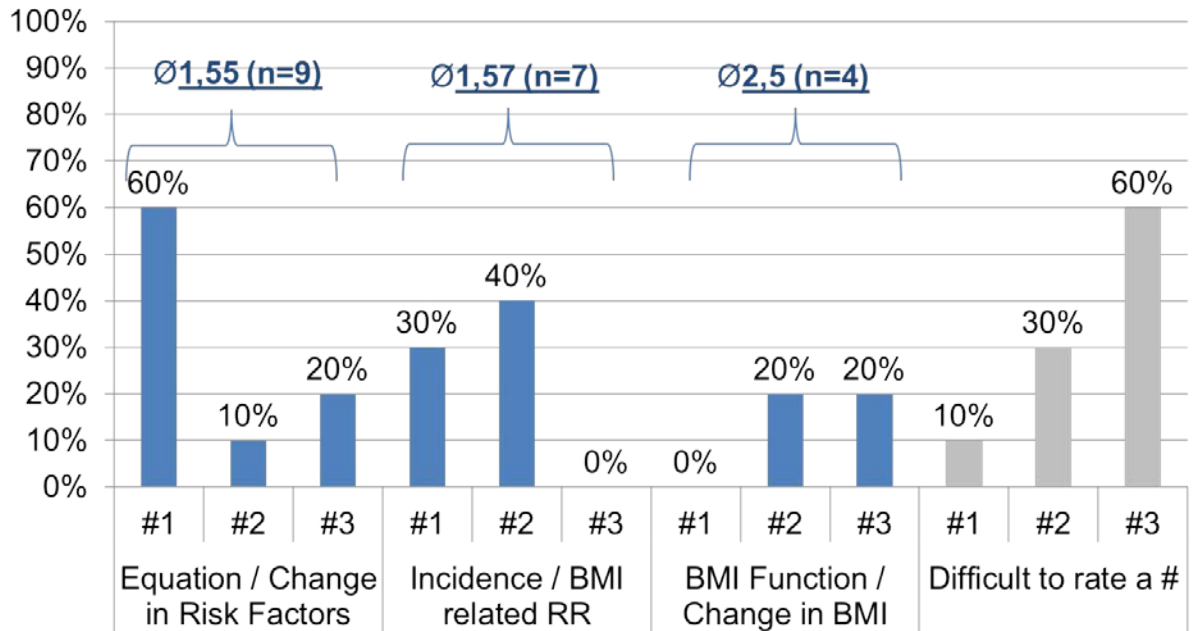


623

624

625

626 **Figure 2: Outcomes of the interview question: Which event simulation**
 627 **approach would you prefer for a health economic obesity model? (Rank 1-3)**



628

629