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# The Role of Diet in the Prevention of Diabetes among Women with Prior Gestational Diabetes: A Systematic Review of Intervention and Observational Studies

## Abstract

**Background** Women with prior gestational diabetes (GDM) have an increased lifetime risk of developing type 2 diabetes mellitus (T2DM). There are no up-to-date systematic reviews analyzing the relationship of diet with risk of developing T2DM following GDM. **Objective** To systematically review the evidence from intervention and observational studies on effects of dietary interventions and associations of dietary intake with T2DM outcomes in women with a GDM history. **Methods** Six electronic databases were searched (Cumulative Index to Nursing and Allied Health Literature, Embase, Medline, Cochrane Central, Proquest, and Scopus) for articles published until May 2019. This review includes intervention and observational studies among women of any age with a history of GDM that reported on the effects of dietary interventions or association of dietary intake (energy, nutrients, foods, dietary patterns) with T2DM, impaired glucose tolerance, impaired fasting glucose, or prediabetes. **Results** The systematic review identified five articles reporting results from four intervention studies, and seven articles reporting results from four observational studies. Findings from intervention studies indicated trends toward beneficial effects of a low-glycemic index diet, a low-carbohydrate diet, and a diet in line with general population dietary guidelines, but studies had unclear or high risk of bias. Findings from two cross-sectional and one prospective study indicated poorer diabetes outcomes for women with higher intakes of branched-chain amino acids, total and heme iron, and a diet relatively low in carbohydrates and high in animal fat and protein, and better outcomes among those consuming diets rich in fruit, vegetables, nuts, fish, and legumes, and low in red and processed meats and sugar-sweetened beverages, after adjustment for confounders, including body mass index. **Conclusions** Findings from observational studies support current dietary guidelines for the prevention of T2DM. Further dietary intervention studies are needed to confirm whether or not dietary modification following a GDM pregnancy reduces women's risk of developing T2DM.

## Keywords

role, gestational, diet, diabetes, systematic, review, intervention, observational, studies, prevention, diabetes, prior, women, among

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**The role of diet in the prevention of diabetes among women with prior gestational diabetes: a systematic review of intervention and observational studies**

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1 **RESEARCH SNAPSHOT:**

2 Research Question: Does diet play a role in the prevention of diabetes among women with a  
3 history of gestational diabetes?

4 Key Findings:

- 5 • This systematic review identified 12 articles published until May 2019, including five  
6 articles reporting on findings from four intervention studies, and seven articles reporting  
7 on findings from four observational studies.
- 8 • Intervention studies generally indicated a trend towards beneficial effects of a low-  
9 glycaemic index diet, a low carbohydrate diet, and a diet in line with general population  
10 guidelines, but had a high risk of bias.
- 11 • Observational studies indicated poorer diabetes outcomes for women with higher intakes  
12 of branched chain amino-acids, total and haem iron and a diet relatively low in  
13 carbohydrates and high in animal fat and protein, and better outcomes for women  
14 consuming diets rich in fruit, vegetables, nuts, fish and legumes, and low in red and  
15 processed meats and sugar-sweetened beverages.

16 **ABSTRACT**

17 **Background:** Women with prior gestational diabetes (GDM) have an increased lifetime risk  
18 of developing type 2 diabetes mellitus (T2DM). There are no up-to-date systematic reviews  
19 analysing the relationship of diet with risk of developing T2DM following GDM.

20 **Objective:** To systematically review the evidence from intervention and observational studies  
21 on effects of dietary interventions and associations of dietary intake with diabetes outcomes  
22 in women with a GDM history.

23 **Methods:** Six electronic databases were searched (CINAHL, EMBASE, Medline, Cochrane  
24 Central, Proquest and Scopus) for articles published until May 2019. This review includes  
25 intervention and observational studies among women of any age with a history of GDM that  
26 reported on the effects of dietary interventions or association of dietary intake (energy,  
27 nutrients, foods, dietary patterns) with T2DM, impaired glucose tolerance, impaired fasting  
28 glucose or pre-diabetes.

29 **Results:** The systematic review identified five articles reporting results from four  
30 intervention studies, and seven articles reporting results from four observational studies.  
31 Findings from intervention studies indicated trends towards beneficial effects of a low-  
32 glycaemic index diet, a low-carbohydrate diet, and a diet in line with general population  
33 dietary guidelines, but studies had unclear or high risk of bias. Findings from two cross-  
34 sectional and one prospective study indicated poorer diabetes outcomes for women with  
35 higher intakes of branched chain amino-acids, total and haem iron and a diet relatively low in  
36 carbohydrates and high in animal fat and protein, and better outcomes among those  
37 consuming diets rich in fruit, vegetables, nuts, fish and legumes, and low in red and  
38 processed meats and sugar-sweetened beverages, after adjustment for confounders including  
39 BMI.

40 **Conclusions:** Findings from observational studies support current dietary guidelines for the  
41 prevention of diabetes. Further dietary intervention studies are needed to confirm if dietary  
42 modification following a GDM pregnancy reduces women's risk of developing T2DM.

## 43 INTRODUCTION

44 Gestational diabetes mellitus (GDM) is defined as “glucose intolerance that begins or is first  
45 diagnosed during pregnancy”.<sup>1</sup> Worldwide, the prevalence of GDM varies from an average of  
46 6% in Europe to 13% in the Middle East and North Africa depending on factors such as  
47 health and lifestyle behaviours, ethnicity and diagnostic criteria,<sup>2</sup> and the incidence is  
48 increasing globally.<sup>3</sup> GDM usually resolves during the postpartum period; however women  
49 with a history of GDM remain a high-risk group for future development of type 2 diabetes  
50 (T2DM). Approximately 50% of women with a history of GDM will go on to develop T2DM  
51 within five to ten years after delivery.<sup>2, 4</sup>

52 T2DM causes numerous complications including retinopathy, kidney failure, cardiovascular  
53 disease and lower limb amputation, increasing an individual’s overall risk of dying  
54 prematurely.<sup>5</sup> The burden of diabetes is enormous worldwide, for example, in 2017 diabetes  
55 was responsible for 4 million deaths and accounted for at least US 727 billion dollars in  
56 health expenditure globally.<sup>6</sup> Interventions that successfully prevent or delay the onset of  
57 T2DM in women with a previous history of GDM have the potential to improve an  
58 individual’s long-term health and reduce the burden of morbidity associated with diabetes.<sup>7</sup>

59 Several systematic reviews and meta-analyses examining the effect of behavioural  
60 interventions, including increasing physical activity and/or improving diet quality, on  
61 diabetes risk in women with prior GDM have previously been published.<sup>7-10</sup> Two reviews that  
62 included meta-analyses highlighted that while ‘intervention’ is generally superior to ‘no  
63 intervention’, effect estimates varied widely between intervention studies.<sup>8, 10</sup> One potential  
64 explanation for this heterogeneity in findings may be that results were pooled from diverse  
65 study types that examined the effects of interventions with goals related to diet modification  
66 alone, physical activity alone, and both diet modification and physical activity.<sup>8, 10</sup> In addition

67 to intervention studies, one review included observational studies and found a limited number  
68 of studies that suggested healthful dietary patterns may be associated with lower risk of  
69 developing T2DM.<sup>7</sup> These previous reviews included studies published until early 2016 (for  
70 intervention studies)<sup>10</sup> or until 2014 (for observational studies).<sup>7</sup> Further systematic reviews  
71 are therefore needed to provide an up-to-date synthesis of the current evidence, incorporating  
72 findings from intervention and observational studies.

73 Interventions that focus on dietary modification alone may have the largest potential to  
74 improve diabetes outcomes in women with a history of GDM. For example, meta-analyses  
75 examining interventions aimed at reducing postpartum weight retention, which may  
76 positively influence diabetes risk, have found that physical activity-only interventions had no  
77 significant effect on postpartum weight loss,<sup>11</sup> while diet-only interventions achieved  
78 significant weight loss.<sup>12</sup> In practice, engagement and compliance with multi-component  
79 interventions, particularly with advice to increase physical activity during the postpartum  
80 period and beyond, may be compromised due to the barriers faced by women at this life-  
81 stage. Barriers such as time constraints related to childcare and returning to work may reduce  
82 the effectiveness of interventions.<sup>13-15</sup>

83 There are currently no up-to-date systematic reviews or meta-analyses that have focussed  
84 specifically on the effects of diet-only interventions or dietary intake on risk of developing  
85 diabetes among women with a previous history of GDM, and that incorporate evidence from  
86 both intervention and observational studies. Therefore, the aim of this research was to  
87 systematically review published intervention and observational studies to identify any effect  
88 of dietary interventions and associations of dietary intake with future diabetes risk for women  
89 with a previous diagnosis of GDM.

## 90 **MATERIALS AND METHODS**

91 The review was registered with PROSPERO International prospective register of systematic  
92 reviews (CRD42018090887) (<http://www.crd.york.ac.uk/PROPSERO>). Methods of the  
93 review were developed using the Cochrane Handbook for Systematic Reviews of  
94 Interventions<sup>16</sup> and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses  
95 (PRISMA) guidelines were used for reporting the methods and outcomes.<sup>17</sup>

#### 96 *Eligibility Criteria*

97 The review included dietary intervention (intervention trials including counselling and/or  
98 food provided by any health professional) or dietary intake (observational) studies targeting  
99 women of any age, with a previous diagnosis of GDM. As the focus of this review included  
100 the prevention of (pre-) diabetes in women with prior GDM, recruitment may have  
101 commenced during pregnancy, however studies were only included if they reported outcomes  
102 and interventions during the postpartum period and beyond. Included interventions were  
103 those promoting a change in dietary intake, rather than nutritional supplement use, or other  
104 lifestyle behaviours e.g. physical activity. Supplement only studies including the provision of  
105 a capsule, pill or tablet containing amino acids, vitamins or minerals were excluded. Trials  
106 where multiple lifestyle interventions were combined, were included if diet was reported as a  
107 separate intervention arm or effect. Dietary intake could be assessed as energy, nutrients,  
108 foods, dietary patterns, or fortified food products. Randomised controlled trials (RCTs),  
109 controlled trials, pre-post studies, cross-sectional, retrospective and prospective studies were  
110 included in the systematic review. When multiple publications were available for the same  
111 study, the publication reporting the greatest number of participants for each outcome was  
112 selected. **Table 1** highlights the PICOS (population, intervention/exposure, comparator,  
113 outcomes, study design) categories for the formulation of the research question.

#### 114 *Search Strategy*

115 The search strategy was developed in consultation with a subject specialist librarian. A  
116 systematic search for publications occurred in November 2017, and an updated search for any  
117 new publications in May 2019. The search was conducted without date limits, using six  
118 electronic databases: CINAHL; EMBASE; Medline; Cochrane Central; Proquest; and  
119 Scopus. The following MeSH terms, words and combinations of words were searched:  
120 (“gestational diabetes” or GDM or “pregnancy diabetes mellitus” or “pregnancy induced  
121 diabetes”); and (history or “medical history” or previous\*); and (diet\* or food\* or  
122 carbohydrate\* or nutrition\* or “glycemic index” or GI); and (“type 2 diabetes” or diabetes or  
123 DM or T2DM or “impaired glucose tolerance” or “prediabetes” or “impaired fasting  
124 glucose”). Keywords were searched as free text in the title, abstract, or topic and combined  
125 using the Boolean operator “AND”. Limits included English language and human subjects.  
126 Details on the search strategy used for Embase are given in **Table 2**. Additional publications  
127 were identified from the reference lists of included articles and from relevant previously  
128 published review articles.

### 129 *Selection Process*

130 All records identified were first assessed for eligibility based on the information contained in  
131 the title and abstract, by two independent reviewers. The full text of all publications that  
132 appeared to meet the eligibility screening (Table 1) was retrieved and subjected to a second  
133 independent assessment for relevance. Any discrepancy in assessment between reviewers was  
134 resolved through discussion.

### 135 *Quality assessment*

136 Selected full texts were assessed for methodological quality by two independent reviewers.  
137 Intervention studies were assessed using the Cochrane risk of bias tool.<sup>18</sup> All intervention

138 studies were coded as low, high or unclear risk in relation to sequence generation, allocation  
139 concealment, blinding of participants and personnel, blinding of outcome assessment,  
140 incomplete outcome data, and selective reporting. The Newcastle-Ottawa Scale (NOS) was  
141 used to assess the quality of nonrandomised studies, including case-control and cohort  
142 studies.<sup>19</sup> A ‘star system’ is used which classifies the studies as good, fair or poor quality in  
143 relation to the selection of study groups, the comparability of groups and the ascertainment of  
144 either the exposure or outcome of interest for case-control and cohort studies respectively.  
145 Any discrepancies were discussed between reviewers and resolved.

#### 146 *Data Extraction*

147 Two independent reviewers extracted relevant data from all included publications. The  
148 extracts were compared, and any difference was verified and resolved through discussion.  
149 The following information was extracted for all studies: study design, country, recruitment  
150 setting, eligibility criteria, population characteristics, duration of follow-up (if applicable),  
151 outcomes examined, outcome assessment and diagnostic criteria, outcome incidence (for  
152 binary outcomes), results, and covariates included in analysis; for intervention studies:  
153 intervention groups, duration, goals, content and compliance; and for observational studies:  
154 exposures examined and exposure assessment.

#### 155 *Data synthesis and Analysis*

156 Due to heterogeneity across studies in dietary factors, study design, and outcomes of interest,  
157 it was not possible to combine study findings in a meta-analysis. Instead, results indicating  
158 significance and direction of the observed associations were qualitatively summarized in  
159 tables for each study. Information on study characteristics was extracted to describe studies  
160 and populations.

## 161 RESULTS

162 The numbers of identified and included studies are shown in **Figure 1**. The database search  
163 (n = 1,201) and screening of bibliographies (n = 3) yielded 1,204 unique articles. After  
164 screening of titles and abstracts, 45 full-text articles were reviewed. Of these, 12 met the  
165 inclusion criteria. Multiple articles were published based on data from a Malaysian dietary  
166 intervention (n = 2)<sup>20, 21</sup> and the Nurses' Health Study II (NHSII) (n = 4).<sup>22-25</sup> Therefore, this  
167 review includes five articles<sup>20, 21, 26-28</sup> reporting on findings from four intervention studies,  
168 and seven articles<sup>22-25, 29-31</sup> reporting on findings from four observational studies.

### 169 Study Characteristics

170 The main characteristics of the included intervention and observational studies are  
171 summarised in **Tables 3 and 4**. Detail on diabetes outcome assessment in each study, and on  
172 intervention goals, content and compliance (intervention studies) and dietary assessment  
173 (observational studies) are described in **Tables 5 and 6**.

#### 174 *Intervention studies*

175 Intervention studies included RCTs conducted in Malaysia (N = 65 to 77)<sup>20, 21</sup> and Australia  
176 (N = 193),<sup>27</sup> and randomised cross-over studies from the USA (N = 17)<sup>26</sup> and Sweden (N =  
177 7)<sup>28</sup> (**Table 3**). Women had a mean baseline age between 31 and 40 years, and mean baseline  
178 BMI between 25kg/m<sup>2</sup> and 28kg/m<sup>2</sup>. The studies from Malaysia, the USA and Sweden  
179 recruited high risk women who were overweight or obese,<sup>20, 21, 26</sup> had prediabetes or a family  
180 history of diabetes,<sup>20, 21</sup> or had a diagnosis of impaired glucose tolerance at 12-months  
181 postpartum.<sup>28</sup> Women were recruited at a median of four months post-GDM pregnancy in the  
182 Malaysian RCT<sup>20, 21</sup> and 1-3 years post-GDM in the Swedish cross-over trial,<sup>28</sup> whereas the  
183 time since GDM pregnancy was not reported in the studies from Australia or the USA.<sup>26, 27</sup>

184 The dietary intervention studies focused on carbohydrate quality (glycaemic index)<sup>20</sup>,  
185 <sup>21, 28</sup> and carbohydrate quantity (percentage of total energy intake),<sup>26</sup> or included general  
186 advice in line with dietary guidelines<sup>27</sup> (Table 5). The Malaysian low-glycaemic index RCT<sup>20</sup>,  
187 <sup>21</sup> included education sessions on restricting energy and fat intake and increasing dietary fibre  
188 intake, with the goal to achieve a 5% to 7% reduction in body weight (for women with BMI  
189 >23 kg/m<sup>2</sup>) or to maintain current weight (BMI <23 kg/m<sup>2</sup>). In the Swedish cross-over trial,  
190 participants were provided with low-glycaemic and high-fibre bread products or high-  
191 glycaemic and low-fibre bread products, with the aim to determine effects on metabolic  
192 parameters.<sup>28</sup> The dietary intervention that focussed on carbohydrate quantity was a cross-  
193 over trial and included an energy-restricted diet where women received nutritional  
194 supplement bars that were identically wrapped and contained 20% protein and either 40% or  
195 55% carbohydrates (i.e. 40% or 25% of energy from fat, respectively) for breakfast, lunch  
196 and snacks, and a meal plan for dinner which comprised a third of total calories. The goal  
197 was to achieve weight loss of 1lb to 2lbs (0.45kg to 0.9kg) per week.<sup>26</sup> The RCT conducted  
198 by Wein and colleagues included provision of a dietary advice sheet ('Target on Healthy  
199 Eating') for intervention and control groups while the intervention group in addition received  
200 phone contact with a dietitian every three months (so possibly 24 times over a six year  
201 follow-up).<sup>27</sup> Interventions were delivered through a combination of one-on-one sessions  
202 with nutritionists, group education sessions, email and short messaging reminders.<sup>20, 21, 26, 27</sup>  
203 Women in the control groups received general dietary recommendations and did not receive  
204 counselling or personal advice. All interventions were focussed on dietary modifications and  
205 did not include specific advice or goals related to physical activity. In line with general  
206 physical activity recommendations, women in the intervention and control groups were  
207 encouraged to engage in moderate physical activity for 30 minutes at least five times a  
208 week<sup>20, 21</sup> or received a reminder of the need for regular exercise.<sup>27</sup> Diabetes related outcomes

209 following the interventions included fasting plasma glucose,<sup>20, 21, 28</sup> serum fasting insulin,<sup>26, 28</sup>  
210 insulin secretion and sensitivity,<sup>28</sup> and T2DM defined using the 1985 WHO guidelines<sup>27</sup>  
211 (Table 5).

### 212 *Observational studies*

213 Observational studies included cross-sectional studies from Canada (N = 281),<sup>30</sup> Korea (N =  
214 381)<sup>29</sup> and Sweden (N = 137),<sup>31</sup> a longitudinal study from the US (the NHSII, N = 3,976 to  
215 4,502),<sup>22-24</sup> and a nested-case control study within the NHSII (N = 347)<sup>25</sup> (Table 4). Women  
216 had a mean baseline age between 34 and 43 years, and a mean baseline BMI between 21  
217 kg/m<sup>2</sup> and 32kg/m<sup>2</sup>. Studies were conducted at different times after GDM pregnancies,  
218 ranging from 6 to 12 weeks<sup>29</sup> to 14 years post-GDM pregnancy.<sup>24</sup>

219 Dietary intake was assessed using validated food frequency questionnaires (FFQs)<sup>22-</sup>  
220 <sup>25, 30, 31</sup> or a 24-hour dietary recall<sup>29</sup> (Table 6). Studies examined overall dietary patterns (low-  
221 carbohydrate dietary pattern,<sup>23</sup> alternate Mediterranean diet (aMED), Dietary Approaches to  
222 Stop Hypertension (DASH) and alternate Healthy Eating Index (aHEI),<sup>24</sup> fruit and vegetable  
223 intake,<sup>30</sup> frequency of meat, fish and vegetarian meals,<sup>31</sup> cooking fat used,<sup>31</sup> macronutrients,<sup>29,</sup>  
224 <sup>31</sup> branched chain amino acids<sup>25</sup> and iron intake<sup>22</sup>). Diabetes related outcomes, including  
225 abnormal glucose tolerance, prediabetes and T2DM, were defined according to the 2017  
226 American Diabetes Association Guidelines,<sup>22-25, 29</sup> the 2013 Canadian Diabetes Association  
227 Guidelines<sup>30</sup> or the 1999 WHO Guidelines<sup>31</sup> (Table 6).

### 228 **Effects of Dietary Interventions on Diabetes Outcomes**

229 Results from dietary intervention studies showed mixed effects on diabetes outcomes (Table  
230 7). Findings from the low-glycaemic index, energy restricted RCT showed no significant  
231 changes in fasting glucose levels after six<sup>21</sup> and 12<sup>20</sup> months in the overall population, but 2-

232 hour plasma glucose had declined significantly more for women in the intervention compared  
233 to the control group after six months.<sup>21</sup> At the 12-month follow-up, fasting plasma glucose  
234 had significantly decreased in the intervention compared with the control group for women  
235 with baseline fasting insulin  $\geq 2 \mu\text{IU}/\text{ml}^{-1}$  but not with baseline fasting insulin  $< 2 \mu\text{IU}/\text{ml}^{-1}$ .<sup>20</sup>  
236 Findings from the cross-over trial providing low-glycaemic and high-fibre bread products  
237 showed no changes in fasting glucose, fasting insulin and insulin sensitivity, but insulin  
238 secretion was lower after consuming the low-glycaemic and high-fibre bread products for  
239 three weeks compared with before.<sup>28</sup> An intervention, with a median follow-up of 51 months  
240 (4 years and 3 months) providing general dietary advice and dietitian contact did not  
241 significantly reduce T2DM risk, although the risk estimate indicated a trend towards a  
242 beneficial effect (relative risk 0.63 (95% CI 0.35, 1.14) relative to the control group who  
243 received no dietitian contact.<sup>27</sup> A cross-over trial comparing different proportions of dietary  
244 energy from carbohydrates for a period of 12 weeks did not show a difference in fasting  
245 insulin levels between dietary interventions.<sup>26</sup>

## 246 **Associations between Dietary Intake and Diabetes Outcomes**

247 Findings from observational studies showed inverse associations between intake of a range of  
248 dietary factors and diabetes outcomes (**Table 8**). Prevalence of prediabetes was 34%<sup>30</sup> and  
249 42%<sup>29</sup> in two cross-sectional studies, and incidence of T2DM ranged from 7% at 6-12 weeks  
250 post-GDM pregnancy<sup>29</sup> to 16% at 14 years post-GDM pregnancy.<sup>22, 23</sup>

251 In terms of overall dietary patterns, results based on the NSHII showed that a diet  
252 relatively low in carbohydrates was associated with higher T2DM risk, especially when fat  
253 and protein intake came largely from animal foods.<sup>23</sup> Findings from the same study also  
254 demonstrated lower T2DM risk with higher *a priori* dietary pattern scores including for the  
255 aMED, DASH and aHEI, after adjustments for key confounders including age, ethnicity,

256 BMI, parity, lifestyle and other dietary factors.<sup>24</sup> While adjustment for BMI attenuated the  
257 risk estimates, statistically significant associations remained when BMI was added to the  
258 model.<sup>23, 24</sup> A nested case-control study within the NHSII found higher intakes of branched  
259 chain amino acids were associated with higher risk of T2DM.<sup>25</sup> The NHSII also showed that  
260 women with the highest compared with the lowest intakes of total iron (dietary and  
261 supplemental), supplemental iron, and dietary haem iron were at higher risk of developing  
262 T2DM, whereas high dietary non-haem iron intake was associated with a lower risk after  
263 adjustment for key confounders.<sup>22</sup> Higher intake of fruit and vegetables was associated with  
264 lower odds of abnormal glucose tolerance in Canadian women based on a cross-sectional  
265 analysis adjusted for age and BMI.<sup>30</sup> Kim and colleagues found in their cross-sectional study  
266 that Korean women who had developed prediabetes or T2DM had higher intakes of total  
267 energy, total and animal protein, and total, animal and monounsaturated fat, after adjustment  
268 for pre-pregnancy BMI.<sup>29</sup> The Swedish cross-sectional study, however, did not find  
269 differences between women with normal glucose tolerance, impaired glucose tolerance and  
270 T2DM in terms of energy and macronutrient intake, after adjustment for key confounders.<sup>31</sup>  
271 This study showed that, compared with women with normal glucose tolerance, women with  
272 impaired glucose tolerance or T2DM had lower proportions of butter used for cooking, and  
273 women with T2DM had higher proportion of margarine used for cooking.<sup>31</sup>

## 274 **Quality Assessment**

### 275 *Intervention studies*

276 Based on the Cochrane risk of bias tool, intervention studies were at high risk of selection  
277 bias (random sequence generation (40% of studies), poor allocation concealment (40% of  
278 studies)) and performance bias (60% of studies) (**Figures 2 and 3**). All five studies had

279 unclear risk of detection and attrition bias, while most studies were scored as low risk of  
280 reporting bias.

### 281 *Observational studies*

282 Quality assessment scores for observational studies ranged from five to seven (out of nine  
283 indicating the lowest degree of bias) based on the NOS-scale (**Table 9**). Main concerns were  
284 1) representativeness of the study sample (no random sample of women in the community),  
285 2) comparability of exposed and unexposed participants based on design or analysis (the two  
286 cross-sectional studies did not adjust for potential key confounding factors, including parity,  
287 family history of diabetes, smoking, physical activity, and other dietary factors) and 3)  
288 adequacy of follow-up rate (studies either did not report on the follow-up rate, or did not  
289 describe the lost population if the follow-up rate was <90%).

290

## 291 **DISCUSSION**

### 292 *Summary of main findings*

293 This systematic review identified a limited number of intervention and observational studies  
294 that examined the role of diet in the prevention of diabetes among women with previous  
295 GDM. Five articles that reported on four small intervention studies found no strong evidence  
296 for a beneficial effect on glucose and insulin levels and on T2DM risk. Four articles that  
297 reported on associations with T2DM risk in a large prospective cohort study showed adverse  
298 associations for higher intakes of branched chain amino-acids, total and haem iron and a low  
299 carbohydrate diet, and protective associations for higher adherence to aMED, AHEI and  
300 DASH diet scores. Cross-sectional findings from one study showed lower intakes of fruit and  
301 vegetables among women with abnormal compared with normal glucose tolerance, while

302 findings from two cross-sectional studies on total energy, protein and fat intake in relation  
303 with prediabetes and impaired glucose tolerance were inconsistent. These observational  
304 findings were adjusted for confounding factors including BMI.

### 305 *Interpretation*

306 The inconclusive findings on the effectiveness of dietary interventions for diabetes  
307 prevention in women with a GDM history are consistent with previous systematic reviews of  
308 RCTs.<sup>7-10</sup> While previous reviews reported on only two diet-alone interventions,<sup>21, 27</sup> this  
309 review identified five articles that reported on four studies.<sup>20, 21, 26-28</sup> These trials were based  
310 on small samples and not powered to detect significant differences.<sup>20, 21, 26-28</sup> These trials also  
311 varied in terms of intervention content (advice focussed on carbohydrate quality,<sup>20, 21, 28</sup>  
312 carbohydrate quantity<sup>26</sup> or overall diet in line with national guidelines<sup>27</sup>), and examined  
313 different outcome measures (glucose<sup>20, 21, 28</sup>, insulin<sup>26, 28</sup> or T2DM<sup>27</sup>). While findings from  
314 intervention studies generally indicated trends towards beneficial effects, the overall quality  
315 of these trials was rated as either unclear or high risk of bias. Based on the current evidence, a  
316 specific dietary intervention for diabetes prevention in women with prior GDM can therefore  
317 not be recommended. Previous systematic reviews have also consistently concluded that  
318 evidence for an effect of combined diet and physical activity interventions is inconclusive,<sup>7-10</sup>  
319 with the exception of strong evidence from the Diabetes Prevention Program (DPP).<sup>32</sup>  
320 Findings from this intensive intervention which focussed on diet and physical activity to  
321 achieve and maintain weight loss of at least 7% of initial body weight showed a greater than  
322 50% reduction in the risk of developing T2DM in women at high risk of T2DM including  
323 women with previous GDM,<sup>32</sup> however, this personalised lifestyle intervention is unlikely to  
324 be feasible for implementation in routine care. As a limited number of studies have examined  
325 diet-alone and physical activity-alone interventions, it remains unclear which diabetes  
326 prevention approach would be most effective for women with a GDM history.<sup>7-10</sup> It has been

327 suggested based on combined diet and physical activity interventions that changes in diet  
328 were more often successfully implemented compared with changes in physical activity.<sup>10</sup>  
329 This may be explained by barriers related to being active and engaging in physical activity  
330 such as lack of time and energy, emotional distress, physical discomfort, and lack of child  
331 care support.<sup>13-15</sup> Further studies should explore the feasibility and acceptability of dietary  
332 interventions for women following childbirth, and future RCTs should accommodate barriers  
333 to improve intervention compliance among new mothers.

334 Findings from observational studies identified in this review showed adverse  
335 associations with diabetes outcomes for higher intakes of branched chain amino-acids,<sup>25</sup> total  
336 and haem iron<sup>22</sup> and a diet relatively low in carbohydrates and high in animal fat and  
337 protein,<sup>23</sup> and beneficial associations for diets rich in fruit, vegetables, nuts, fish and legumes,  
338 and low in red and processed meats and sugar-sweetened beverages.<sup>24, 30</sup> These results  
339 support dietary recommendations for the general population<sup>33, 34</sup> and for diabetes  
340 prevention.<sup>35</sup> Findings from observational studies were rated to be of medium to high quality  
341 (score of 5-7 out of 9), and observed associations were based on analyses adjusted for BMI<sup>22-</sup>  
342 <sup>25, 29, 30</sup> and physical activity.<sup>22-25</sup> These findings therefore suggest that, while BMI partly  
343 attenuated the associations,<sup>23, 24</sup> consuming a healthy diet may help reduce T2DM risk  
344 independent of BMI.

#### 345 *Implications for practice and research*

346 While there is strong evidence that the majority of T2DM cases in the general population can  
347 be prevented through a healthy lifestyle,<sup>35, 36</sup> lifestyle interventions targeting women with  
348 prior GDM have only recently emerged. With the growing rates of GDM, T2DM and its  
349 associated health and economic burden worldwide,<sup>5, 6</sup> interventions that successfully prevent  
350 T2DM in this high-risk group are critical. Postnatal screening following a GDM pregnancy

351 represents an important opportunity for clinicians and health professionals to educate, counsel  
352 and engage women in prevention programs that may reduce their risk of progressing to  
353 diabetes. However, the uptake of postnatal follow-up by women with GDM is poor with less  
354 than 25% of women returning for a postnatal glucose tolerance test, mainly due to lack of  
355 time and awareness of risks.<sup>37</sup> Previous studies have shown poor diet quality and low  
356 adherence to dietary recommendations among women with a GDM history,<sup>38,39</sup> and a  
357 significant knowledge gap on future health risks and the recommended lifestyle  
358 modifications.<sup>39</sup> While a causal effect of diet on T2DM following GDM has not been  
359 established, and further research is needed to develop effective postnatal preventive  
360 interventions, it seems prudent to inform women about their future risk of developing T2DM  
361 and the potential benefit of a healthy diet.

#### 362 *Strengths and limitations*

363 Strengths of this systematic review include the focus on dietary intake, which has not been  
364 distinguished from effects of physical activity in previous systematic reviews. Moreover, this  
365 review includes all study designs (both intervention and observational), providing a more  
366 comprehensive overview of the current evidence. This review is restricted, however, by the  
367 lack of high-quality dietary intervention studies. Further evidence from postpartum dietary  
368 RCTs that follow women for a longer period of time are needed to confirm findings from  
369 observational studies. Current observational findings suggest that future RCTs should not  
370 only focus on carbohydrate quality and quantity, but also on other aspects of the overall diet  
371 including increasing intake of fruit, vegetables and other plant-based foods, and reducing  
372 animal products such as red and processed meats.

#### 373 **CONCLUSIONS**

374 This systematic review indicates a lack of high quality dietary RCT evidence for the  
375 prevention of diabetes among women with prior GDM. Findings from observational studies  
376 have shown beneficial associations of several nutrients, foods and dietary patterns in line with  
377 current dietary recommendations and suggest diet may play a role in the prevention of  
378 diabetes among women with prior GDM. Further large prospective dietary RCTs that are  
379 acceptable to new mothers are needed to confirm if dietary modifications in line with healthy  
380 eating guidelines reduce the risk of T2DM among women following a GDM pregnancy.

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## FIGURE LEGEND

**Figure 1.** Flow diagram of the literature search and screening process for a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

**Figure 2.** Risk of bias for each individual intervention study included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes. Risk of bias was assessed using the Cochrane risk of bias tool (Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0. The Cochrane Collaboration, 2011. Available from [www.cochrane-handbook.org](http://www.cochrane-handbook.org)).

**Figure 3.** Risk of bias summary for five intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes. Risk of bias was assessed using the Cochrane risk of bias tool (Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0. The Cochrane Collaboration, 2011. Available from [www.cochrane-handbook.org](http://www.cochrane-handbook.org)).

## **TABLES**

**Table 1.** PICOS categories for formulation of the research question for a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

**Table 2.** Search strategy for Embase in a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

**Table 3.** Characteristics of intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

**Table 4.** Characteristics of observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

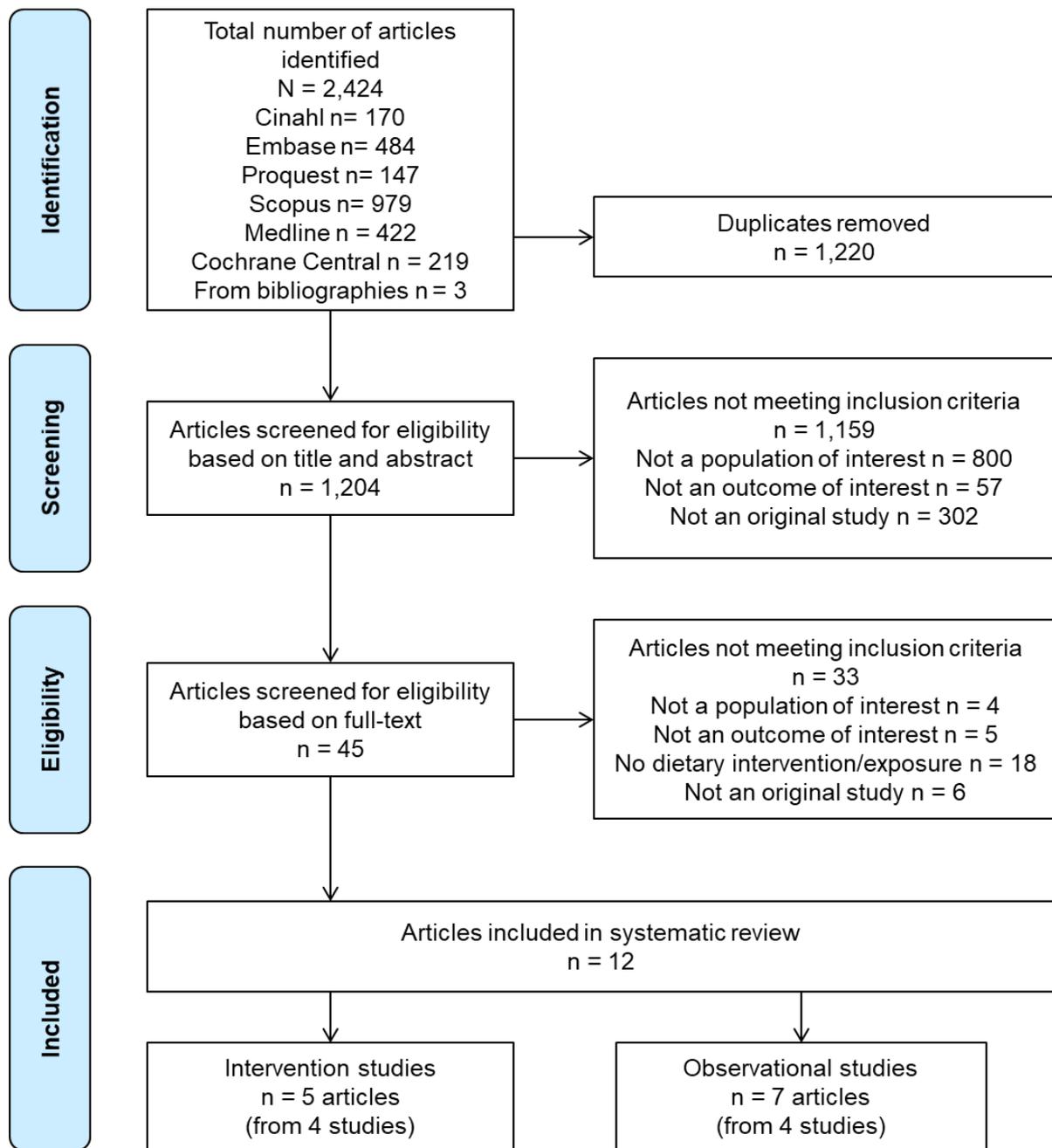
**Table 5.** Intervention goals, content and compliance, and diabetes outcome assessment in intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

**Table 6.** Diet and diabetes outcome assessment in observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

**Table 7.** Results from intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

**Table 8.** Results from observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

**Table 9.** Quality assessment for each individual observational study included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

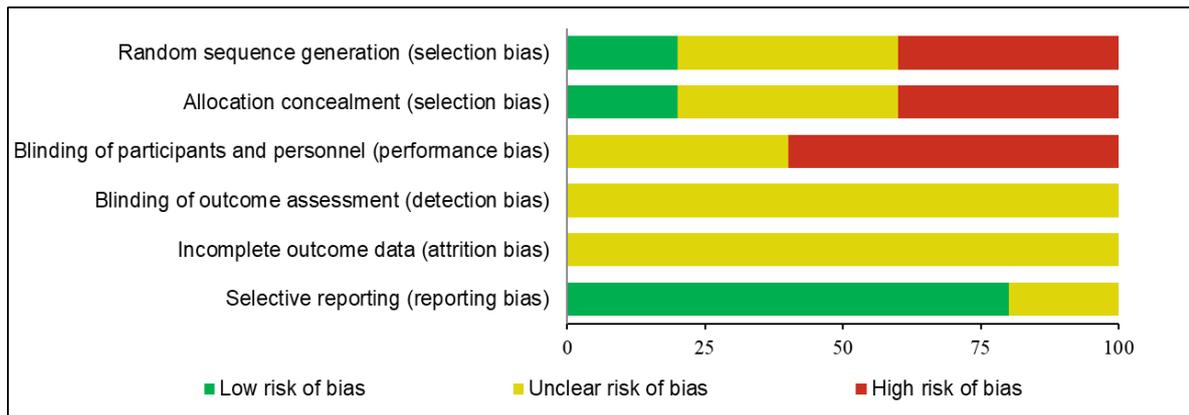


**Figure 1.** Flow diagram of the literature search and screening process for a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)
Ghani RA, 2014 <sup>20</sup>	-	-	-	?	?	+
Shyam S, 2013 <sup>21</sup>	+	+	-	?	?	+
Östman E, 2006 <sup>28</sup>	-	-	?	?	?	?
Wein P, 1999 <sup>27</sup>	?	?	-	?	?	+
Peterson CM, 1995 <sup>26</sup>	?	?	?	?	?	+

 Low risk   
 Unclear risk   
 High risk

**[online] Figure 2.** Risk of bias for each individual intervention study included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes. Risk of bias was assessed using the Cochrane risk of bias tool (Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0. The Cochrane Collaboration, 2011. Available from [www.cochrane-handbook.org](http://www.cochrane-handbook.org)).



**[online] Figure 3.** Risk of bias summary for five intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes. Risk of bias was assessed using the Cochrane risk of bias tool (Higgins JPT, Green S (editors). *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.1.0. The Cochrane Collaboration, 2011. Available from [www.cochrane-handbook.org](http://www.cochrane-handbook.org)).

**Table 1.** PICOS categories for formulation of the research question for a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

Population	Women (postpartum onwards) with a history of gestational diabetes
Intervention/ exposure	Dietary intervention (intervention studies), or dietary intake (energy, nutrients, foods, dietary patterns) (observational studies)
Comparison	Control group (e.g. standard care as part of randomised controlled trial) or no control group (e.g. pre-post intervention study), not applicable for observational studies
Outcome	Type 2 diabetes, impaired glucose tolerance, impaired fasting glucose, or prediabetes
Study designs	Intervention studies (no restrictions on randomisation and inclusion of a control group), observational studies (cross-sectional, retrospective and prospective studies)

**[online] Table 2.** Search strategy for Embase in a systematic review on effects of dietary interventions and associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

Search #	Search term <sup>a</sup>
1	gestational diabetes:ti,ab
2	GDM:ti,ab
3	pregnancy diabetes mellitus:ti,ab
4	pregnancy induced diabetes:ti,ab
5	1 or 2 or 3 or 4
6	history:ti,ab
7	medical history:ti,ab
8	previous*:ti,ab
9	6 or 7 or 8
10	diet*:ti,ab
11	food*:ti,ab
12	carbohydrate*:ti,ab
13	nutrition*:ti,ab
14	glycemic index:ti,ab
15	GI:ti,ab
16	10 or 11 or 12 or 13 or 14 or 15
17	type 2 diabetes:ti,ab
18	diabetes:ti,ab
19	DM:ti,ab
20	T2DM:ti,ab
21	impaired glucose tolerance:ti,ab
22	prediabetes:ti,ab
23	impaired fasting glucose:ti,ab
24	17 or 18 or 19 or 20 or 21 or 22 or 23
25	5 and 9 and 16 and 24

<sup>a</sup> Restricted to English language and human populations. Similar searches were conducted in all other databases.

**Table 3.** Characteristics of intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

First author, year	Country	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
Ghani RA, 2014 <sup>20</sup>	Malaysia	Randomised controlled trial N = 77 (n = 39 in low glycaemic index intervention; n = 38 in control group)	Included: women with a history of gestational diabetes aged 20-40 years, without a current diagnosis of diabetes, at risk of type 2 diabetes (at least one of the four following risk factors: BMI > 23 kg/m <sup>2</sup> , or waist circumference > 80 cm, or impaired glucose tolerance or impaired	Women were recruited from the endocrine clinic of a tertiary hospital.	Time since gestational diabetes pregnancy: median 4 months Age at baseline (mean): 31 years Previous births (mean): 2	(1) Low glycaemic index intervention, (2) Control group. Intervention duration not reported. Women were followed for 12 months post-intervention.	Fasting plasma glucose

First author, year	Country	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
			fasting glucose, or a family history of type 2 diabetes). Excluded: women with a diagnosis of type 2 diabetes, or presence of other health complications and usage of drugs affecting body weight and glucose control.		BMI (mean): 26 kg/m <sup>2</sup>		
Shyam S, 2013 <sup>21</sup>	Malaysia	Randomised controlled trial	Included: women with a history of gestational diabetes aged 20-40 years,	Women were recruited from the	Time since gestational diabetes	(1) Low glycaemic index	Fasting plasma glucose,

First author, year	Countr y	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
		N = 65 women (n = 34 in low glycaemic index intervention; n = 31 in the control group)	without a current diagnosis of diabetes, at risk of type 2 diabetes (at least one of the four following risk factors: BMI>23 kg/m <sup>2</sup> , or waist circumference>80 cm, or impaired glucose tolerance or impaired fasting glucose, or a family history of type 2 diabetes). Excluded: women with a diagnosis of type 2 diabetes, or	endocrine clinic of a tertiary hospital.	pregnancy: median 4 months Age at baseline (mean): 31 years Previous births (mean): 2 BMI (mean): 26 kg/m <sup>2</sup>	intervention, (2) Control group. Intervention duration not reported. Women were followed for 6 months post- intervention.	2-hours glucose

First author, year	Country	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
			presence of other health complications and usage of drugs affecting body weight and glucose control.				
Östman E, 2006 <sup>28</sup>	Sweden	Randomized cross-over study N = 7	Included: Women with a history of gestational diabetes 1-3 years prior the study, who had a diagnosis of impaired glucose tolerance at 12-months postpartum.	Not reported	Age at baseline (mean): 32 years BMI (mean): 28 kg/m <sup>2</sup>	(1) Dietary intervention 1 (2) Dietary intervention 2. Participants were crossed over after a 3-week	Insulin secretion, insulin sensitivity, fasting glucose and insulin

First author, year	Countr y	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
Wein P, 1999 <sup>27</sup>	Australi a	Randomised controlled trial	Included: Women with a history of gestational diabetes who had a	Women were recruited from the Mercy Hospital for	Age at baseline (mean): 38-40 years	intervention and 3-week washout period (total duration is 9 weeks). No control group. No post- intervention follow-up. (1) Lifestyle intervention, (2) Control group.	Type 2 diabetes

First author, year	Countr y	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
		N = 193 (n = 97 in intervention group; n = 96 in control group)	diagnosis of impaired glucose tolerance and who could communicate directly or through translation. No exclusion criteria reported.	Women in Melbourne between November 1989 and July 1991	Previous births (mean): 3 BMI (mean): 25- 26 kg/m <sup>2</sup>	Intervention duration not reported. Women were followed for up to 6 years (median follow- up of 51 months (4 years and 3 months)) post- intervention.	

First author, year	Country	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
Peterson CM, 1995 <sup>26</sup>	USA	Randomised cross-over study N = 17	Included: Women with a history of gestational diabetes who were overweight or obese, 1-4 years postpartum, not nursing, and aged 21-50 years. Excluded: women with any medical condition for which a calorie-restricted diet may be harmful including pregnancy or planned pregnancy during the trial,	Not reported	Age at baseline (mean): 34-38 years Previous births (mean): 3 BMI (mean): 25- 26 kg/m <sup>2</sup>	(1) Dietary intervention 1 (2) Dietary intervention 2. Participants were crossed over after a 6- week intervention (total duration is 12 weeks). No control group.	Serum fasting insulin

First author, year	Country	Study design, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Intervention groups, duration and follow-up	Outcome
			hypertension, diuretics use, thyroid disease, or diabetes.			No post- intervention follow-up.	

<sup>a</sup> Participant characteristics extracted were time since gestational diabetes pregnancy, age, ethnicity, parity and BMI, if reported. Characteristics are reported as a range across categories when no descriptive statistics were presented for the overall study population.

**Table 4.** Characteristics of observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
Tobias DK, 2018 <sup>25</sup>	USA	Prospective nested case – control study (Nurses’ Health Study II, as part of the Diabetes and Women’s Health Study) N = 172 diabetes cases	Included: women with prevalent gestational diabetes before 1991, or incident gestational diabetes between 1991 and 2001, and with a plasma sample available from the biospecimen collection.	In 1989, nurses aged 25-42 years in California, Connecticut, Indiana, Iowa, Kentucky, Massachusetts, Michigan, Missouri, New York, North Carolina, Ohio, Pennsylvania, South Carolina, and Texas	Years since first gestational diabetes pregnancy (mean): 12 years (cases) and 3 years (controls) Age at baseline blood draw (mean): 43 years Ethnicity: 91% (cases) 94% (controls) Caucasian	Total and subtypes of branched-chain amino acids: isoleucine, leucine and valine	Type 2 diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
		and 175 age-matched controls	Excluded: women with cancer, missing dietary data, or missing metabolite data.	were recruited through invitations.	BMI (mean): 31.6 (cases) and 25.5 (controls) kg/m <sup>2</sup>		
Mercier R, 2018 <sup>30</sup>	Canada	Cross-sectional No follow-up N = 281	Included: women aged ≥18 with a diagnosis of gestational diabetes between 2003 and 2013. Excluded: women pregnant during the study or	Women were recruited since 2009 through a provincial health plan registry and through medical records from two major hospitals	Time since gestational diabetes pregnancy (mean): 6 years Age at baseline (mean): 37 years Previous births (mean): 2	Fruit and vegetable intake	Abnormal glucose tolerance, including impaired fasting glucose, impaired glucose

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
			with diabetes before pregnancy.	within a neonatal care unit in Quebec City.	BMI (mean): 26-29 kg/m <sup>2</sup>		tolerance, prediabetes and type 2 diabetes
Andersson-Hall U, 2018 <sup>31</sup>	Sweden	Cross-sectional No follow-up N = 137	Included: women diagnosed with gestational diabetes between 2005 and 2009.	Women were recruited in the Gothenburg area through a telephone interview.	Time since gestational diabetes pregnancy (mean): 6 years Age at baseline (mean): 38-41 years Ethnicity: 33-53% Scandinavian BMI (mean): 26-30 kg/m <sup>2</sup>	Intake of energy, carbohydrates, protein and fat, frequency of meat, fish and vegetarian meals, and type of cooking fat used	Impaired glucose tolerance and type 2 diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
						(proportion of butter, margarine and vegetable oil)	
Bao W, 2016(a) <sup>22</sup>	USA	Prospective cohort (Nurses' Health Study II, as part of the Diabetes and Women's Health Study)	Included: women with prevalent gestational diabetes before 1991, or incident gestational diabetes between 1991 and 2001. Excluded: women with chronic disease	In 1989, nurses aged 25-42 years in California, Connecticut, Indiana, Iowa, Kentucky, Massachusetts, Michigan, Missouri, New York, North Carolina, Ohio,	Age at baseline (mean): 38 years Ethnicity: 91-93% Caucasian BMI (mean): 26-27 kg/m <sup>2</sup>	Dietary total iron, heme iron, non-heme iron, and supplemental iron (from multivitamin and specific	Type 2 diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
		Followed-up from 1991 to 2001, at 4-yearly intervals N = 3,976	(type 2 diabetes, cardiovascular disease, or cancer) before gestational diabetes pregnancy or before return of first post-gestational diabetes food frequency questionnaire, multiple pregnancy, no post-gestational diabetes food	Pennsylvania, South Carolina, and Texas were recruited through invitations.		iron supplements)	

First author, year	Country	Study design, follow-up, n women	Eligibility criteria frequency questionnaire.	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
Bao W, 2016(b) <sup>23</sup>	USA	Prospective cohort (Nurses' Health Study II, as part of the Diabetes and Women's Health Study)	Included: women with prevalent gestational diabetes before 1991, or incident gestational diabetes between 1991 and 2001. Excluded: women with chronic disease	In 1989, nurses aged 25-42 years in California, Connecticut, Indiana, Iowa, Kentucky, Massachusetts, Michigan, Missouri, New York, North Carolina, Ohio,	Age at baseline (mean): 38 years Ethnicity: 91-93% Caucasian >1 previous birth: 80-84% BMI (mean): 25-29 kg/m <sup>2</sup>	Overall (based on total carbohydrate, protein and fat intake), animal (based on total carbohydrate, animal protein and animal fat	Type 2 diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
		Followed-up from 1991 to 2001, at 4-yearly intervals N = 4,502	(type 2 diabetes, cardiovascular disease, or cancer) before gestational diabetes pregnancy or before return of first post-gestational diabetes food frequency questionnaire, multiple pregnancy, no post-gestational diabetes food	Pennsylvania, South Carolina, and Texas were recruited through invitations.		intake) and vegetable low-carbohydrate dietary pattern (based on total carbohydrate, vegetable protein and vegetable fat)	

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
			frequency questionnaire.				
Tobias DK, 2012 <sup>24</sup>	USA	Prospective cohort (Nurses' Health Study II, as part of the Diabetes and Women's Health Study)	Included: women with prevalent gestational diabetes before 1991, or incident gestational diabetes between 1991 and 2001. Excluded: women with chronic disease	In 1989, nurses aged 25-42 years in California, Connecticut, Indiana, Iowa, Kentucky, Massachusetts, Michigan, Missouri, New York, North Carolina, Ohio,	Time from gestational diabetes pregnancy to type 2 diabetes diagnosis (mean): 13.8 years Age at baseline (mean): 38 years Ethnicity: 91-93% Caucasian	Three a priori dietary pattern scores: the alternate Mediterranean diet (aMED), Dietary Approaches to Stop	Type 2 diabetes

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
		Followed-up from 1991 to 2001, at 4-yearly intervals N = 4,413	(type 2 diabetes, cardiovascular disease, or cancer) before gestational diabetes pregnancy or before return of first post-gestational diabetes food frequency questionnaire, multiple pregnancy, no post-gestational diabetes food	Pennsylvania, South Carolina, and Texas were recruited through invitations.	BMI (mean): 21-22 kg/m <sup>2</sup>	Hypertension (DASH) and alternate Healthy Eating Index (aHEI)	

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
			frequency questionnaire, more than 70 food frequency questionnaire items left blank or unrealistic energy intake (<500 kcal/day or >3500 kcal/day).				
Kim S-H, 2011 <sup>29</sup>	Korea	Cross-sectional No follow-up N = 381	Included: women with a recent gestational diabetes pregnancy and	Between June 2006 and March 2009, pregnant women visiting the Cheil	Time since gestational diabetes pregnancy: 6-12 weeks	Total energy, carbohydrates, total, animal and plant	Type 2 diabetes and prediabetes (impaired glucose

First author, year	Country	Study design, follow-up, n women	Eligibility criteria	Recruitment setting	Participant characteristics <sup>a</sup>	Exposure(s)	Outcome(s)
			normal glucose levels in early pregnancy. Excluded: women with b-cell-specific autoantibodies including glutamic acid decarboxylase antibodies (GADA).	General Hospital (Seoul, Korea) were recruited.	Age at baseline (mean): 34-35 years No previous birth: 40-46% Pre-pregnancy BMI (mean): 23-25 kg/m <sup>2</sup>	protein, total, animal, plant, saturated, mono- and polyunsaturated fat, omega 3 and 6 fatty acids, dietary fibre	tolerance and/or impaired fasting glucose)

<sup>a</sup> Participant characteristics extracted were time since gestational diabetes pregnancy, age, ethnicity, parity and BMI, if reported. Characteristics are reported as a range across categories when no descriptive statistics were presented for the overall study population.

**[online] Table 5.** Intervention goals, content and compliance, and diabetes outcome assessment in intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

First author, year	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
Ghani RA, 2014 <sup>20</sup>	(1) Low glycaemic index intervention group: women received conventional healthy dietary recommendations including nutrition education at baseline through a structured one-on-one session with a research nutritionist, sample menus, and take-home reference booklets. Education sessions emphasised restriction of energy and fat intake and encouraged increases in dietary fibre intake. The goal was to achieve a 5-7% reduction in body weight if BMI > 23 kg/m <sup>2</sup> or to maintain current weight if BMI < 23 kg/m <sup>2</sup> . Energy requirements were calculated, and women with BMI > 23 kg/m <sup>2</sup> had a 500kcal energy	Compliance was monitored through assessments of dietary intake (3-day dietary records at baseline, 3 and 6 months), and nutrition knowledge assessment pertaining to the group-specific concepts. After six months, mean percentage of calories from carbohydrates, and glycaemic index and	Plasma glucose was measured on blood samples drawn in the fasting state (12 hours).  Diagnostic criteria: not applicable (continuous outcome).

First author,	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
year	<p>restricted diet, capped at 1800kcal/day. Women were followed-up quarterly, and received fortnightly reminders using email or short messaging services reinforcing concepts of healthy living and motivating women to comply with the intervention. Women were also encouraged to engage in moderate physical activity for 30 minutes at least five times a week.</p> <p>In addition to conventional healthy dietary recommendations, women received low glycaemic index education including advice on how to choose low glycaemic index options in place of high glycaemic index staples like bread, rice and breakfast cereal. Women were</p>	<p>glycemic load were significantly lower in the intervention compared to the control group. Fibre intake was significantly higher in the intervention group.</p>	

First author,	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
year	<p>asked to restrict rice intake to once/day and include 1 low glycaemic index food at each meal.</p> <p>(2) Control group: women received conventional healthy dietary recommendations only.</p>	See above	<p>Plasma glucose was measured on blood samples drawn in the fasting state (12 hours). 2-hour post load blood glucose was measured after a 75 g oral glucose tolerance test.</p> <p>Diagnostic criteria: not applicable (continuous outcome).</p>
Shyam S, 2013 <sup>21</sup>	See above	See above	

First author,	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
Östman E, 2006 <sup>28</sup>	<p>(1) Low glycaemic index and high fibre bread products.</p> <p>(2) High glycaemic index and low fibre bread products.</p> <p>Participants were asked to eat seven slices of low glycaemic index /high fibre or high glycaemic index /low fibre bread across three meals per day. Participants were also supplied with butter, cheese and ham to be used on the bread in amounts according to written instructions. Apart from the bread meals, participants were asked to maintain their habitual diet throughout the study, but were not allowed to eat any other type of bread.</p>	Not reported	<p>Fasting glucose and insulin were measured on blood samples drawn in the fasting state. Insulin secretion and insulin sensitivity were assessed based on an intravenous glucose tolerance test followed by a euglycaemic-hyperinsulinaemic clamp.</p> <p>Diagnostic criteria: not applicable (continuous outcomes).</p>

First author, year	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
Wein P, 1999 <sup>27</sup>	<p>(1) Lifestyle intervention: women were given dietary questionnaires and a standard dietary advice sheet ('Target on Healthy Eating') and in addition had 3-monthly phone contact with a dietitian.</p> <p>(2) Control group: women were given dietary questionnaires and a standard dietary advice sheet ('Target on Healthy Eating') only.</p> <p>Both groups received a reminder of the need for regular exercise.</p>	<p>A diet score (based on the fat, residue and sugar content of the diet assesses through diet history) was assessed and not significantly different at trial entry between control and intervention groups, and were also not different at final assessment (diet improved in both groups).</p>	<p>Type 2 diabetes was assessed using an oral glucose tolerance test after an overnight fast.</p> <p>Diagnostic criteria: World Health Organisation guidelines 1985.</p>
Peterson CM, 1995 <sup>26</sup>	<p>(1) Dietary intervention 1: 40% carbohydrates</p> <p>(2) Dietary intervention 2: 55% carbohydrates</p>	<p>Women were instructed to maintain a daily diary wherein they recorded all</p>	<p>Serum fasting insulin was assessed at baseline and after 6 and 12 weeks.</p>

First author,	Intervention goals and content	Intervention compliance	Outcome assessment and criteria
year	<p>Women received nutritional supplement bars for breakfast, lunch, and snacks. These were identically wrapped, each contained 180 kcal with 20% protein, and either 40% or 55% carbohydrates.</p> <p>A meal plan for dinner was provided.</p> <p>The daily caloric prescription consisted of 16.5 kcal/kg bodyweight at entry into the trial with a mean caloric consumption of 1500 kcal/24 hour for both groups.</p> <p>The anticipated weight loss was calculated at 1 to 2 lbs/week.</p>	<p>foods consumed by time of day. These were reviewed every one or two weeks. All participants showed significant weight loss in the first 6 weeks regardless of treatment assignment. Weight loss attenuated during the second 6 weeks of the study also regardless of treatment assignment. The diet score at enrolment did not differ between the intervention and</p>	<p>Diagnostic criteria: not applicable (continuous outcome).</p>

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First author,

year

Intervention goals and content

Intervention compliance

Outcome assessment and criteria

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control groups respectively

(3.1 and 3.2, respectively).

There was improvement in

diet scores, but this was

comparable in the

intervention and control

groups (0.64 and 0.56,

p=0.32).

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**[online] Table 6.** Diet and diabetes outcome assessment in observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

First author, year	Exposure assessment	Outcome assessment and criteria
Mercier R, 2018 <sup>30</sup>	Validated 91-item food frequency questionnaire reflecting past month consumption	Glycemic measurements were obtained during each woman's visit at the clinical investigation unit.
	Administered by a registered dietitian during a clinical visit	Diagnostic criteria: Canadian Diabetes Association Guidelines 2013 Impaired fasting glucose: fasting glucose 6.1 to 6.9 mmol/L and normal 2-hour post-oral glucose tolerance test glucose < 7.8 mmol/L Impaired glucose tolerance: normal fasting glucose < 6.1 mmol/L and 2-hour post-oral glucose tolerance test glucose 7.8 to 11.0 mmol/L Prediabetes: impaired fasting glucose and/or impaired glucose tolerance and/or A1C between 6.0 and 6.4% Type 2 diabetes: fasting glucose $\geq$ 7.0 mmol/L and/or 2-hour post-oral glucose tolerance test glucose $\geq$ 11.1 mmol/L and/or A1C value $\geq$ 6.5%

First author, year	Exposure assessment	Outcome assessment and criteria
Bao W, 2016(a) <sup>22</sup>	Validated semi- quantitative food frequency questionnaire.	Type 2 diabetes diagnosis was self-reported. A validation study showed 94% of self-reported cases were confirmed by medical records.
Bao W, 2016(b) <sup>23</sup>	Administered as part of survey questionnaire	Diagnostic criteria: American Diabetes Association Guidelines 1997 Type 2 diabetes: 1) one or more classic symptoms (excessive thirst, polyuria, unintentional weight loss, or hunger) plus elevated glucose concentrations (fasting plasma glucose $\geq 7.0$ mmol/L or random plasma glucose $\geq 11.1$ mmol/L); 2) no symptoms reported but $\geq 2$ elevated plasma glucose concentrations on more than one occasion (fasting glucose $\geq 7.0$ mmol/L; random glucose $\geq 11.1$ mmol/L, or 2-hour post-oral glucose tolerance test glucose $\geq 11.1$ mmol/L); or 3) treatment with insulin or an oral hypoglycemic agent. Before 1998, fasting plasma glucose $\geq 7.8$ mmol/L was used instead of $\geq 7.0$ mmol/L according to the criteria of the National Diabetes Data Group.
Tobias DK, 2012 <sup>24</sup> and 2018 <sup>25</sup>	Validated semi- quantitative food frequency questionnaire.	Blood samples were taken at the study visit 6 years after pregnancy. Diagnostic criteria: 1999 World Health Organisation guidelines Type 2 diabetes: fasting glucose $\geq 7.0$ mmol/L or 2-hour post-oral glucose tolerance test glucose $\geq 11.1$ mmol/L
Andersson- Hall U, 2018 <sup>31</sup>		

First author, year	Exposure assessment	Outcome assessment and criteria
Kim S-H, 2011 <sup>29</sup>	Completed during study visit. 24-hour recall Administered by a skilled dietitian	Impaired glucose tolerance: fasting glucose 6.1 to 6.9 mmol/L and/or 2-hour post-oral glucose tolerance test glucose 7.8 to 11.0 mmol/L Type 2 diabetes and prediabetes were assessed at a clinical visit at 6-12 weeks postpartum Diagnostic criteria: American Diabetes Association Guidelines 1997 Type 2 diabetes: fasting glucose >7.0 mmol/L or 2-hour glucose >11.1 mmol/L Impaired glucose tolerance: fasting glucose <6.1 mmol/L and 2-hour glucose 7.8 to 11.0 mmol/L Impaired fasting glucose: fasting glucose 6.1 to 6.9 mmol/L inclusive with 2-hour glucose <7.8 mmol/L

**Table 7.** Results from intervention studies included in a systematic review on effects of dietary interventions on diabetes outcomes in women with a history of gestational diabetes

First author, year	Outcome incidence	Results	Covariates included in analysis
Ghani RA, 2014 <sup>20</sup>	Not applicable (continuous outcome)	<p>Changes in fasting plasma glucose (mean <math>\pm</math> standard deviation):</p> <p>Women with fasting insulin <math>&lt;2 \mu\text{U}/\text{ml}^{-1}</math>:</p> <ul style="list-style-type: none"> <li>- Intervention group: <math>0.48 \pm 1.2 \text{ mmol/L}</math></li> <li>- Control group: <math>0.18 \pm 0.32 \text{ mmol/L}</math> (<math>p = 0.16</math> for intervention vs control)</li> </ul> <p>Women with fasting insulin <math>\geq 2 \mu\text{U}/\text{ml}^{-1}</math>:</p> <ul style="list-style-type: none"> <li>- Intervention group: <math>-0.12 \pm 0.27 \text{ mmol/L}</math></li> <li>- Control group: <math>0.17 \pm 0.32 \text{ mmol/L}</math> (<math>p = 0.03</math> for intervention vs control)</li> </ul>	None/not reported
Shyam S, 2013 <sup>21</sup>	Not applicable (continuous outcome)	<p>Overall:</p> <p>Changes in fasting plasma glucose (mean <math>\pm</math> standard deviation):</p> <ul style="list-style-type: none"> <li>- Intervention group: <math>-0.2 \pm 0.6 \text{ mmol/L}</math></li> </ul>	Glycaemic load and carbohydrate intake

First author,	Outcome	Results	Covariates included in analysis
year	incidence	<p data-bbox="696 347 1570 451">- Control group: <math>0.1 \pm 0.6</math> mmol/L (p = &gt;0.05 for intervention vs control)</p> <p data-bbox="651 496 1099 523">Changes in 2-hour plasma glucose:</p> <p data-bbox="696 571 1234 598">- Intervention group: <math>-0.2 \pm 2.8</math> mmol/L</p> <p data-bbox="696 646 1570 746">- Control group: <math>0.8 \pm 2.0</math> mmol/L (p = 0.03 for intervention vs control)</p> <p data-bbox="651 866 1144 893">Women with baseline normoglycemia:</p> <p data-bbox="651 941 1469 968">Changes in fasting plasma glucose (mean <math>\pm</math> standard deviation):</p> <p data-bbox="696 1016 1256 1043">- Intervention group: <math>0.31 \pm 0.53</math> mmol/L</p> <p data-bbox="696 1091 1570 1192">- Control group: <math>0.15 \pm 0.41</math> mmol/L (p = 0.29 for intervention vs control)</p> <p data-bbox="651 1240 1469 1267">Changes in 2-hour plasma glucose (mean <math>\pm</math> standard deviation):</p> <p data-bbox="696 1315 1256 1342">- Intervention group: <math>0.24 \pm 2.04</math> mmol/L</p>	

First author, year	Outcome incidence	Results	Covariates included in analysis
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- Control group:  $0.88 \pm 1.2$  mmol/L (p = 0.22 for intervention vs control)

Women with baseline dysglycaemia:

Changes in fasting plasma glucose (mean  $\pm$  standard deviation):

- Intervention group:  $0.49 \pm 1.7$  mmol/L
- Control group:  $0.04 \pm 0.45$  mmol/L (p = 0.51 for intervention vs control)

Changes in 2h plasma glucose (mean  $\pm$  standard deviation):

- Intervention group:  $-0.49 \pm 4.05$  mmol/L
- Control group:  $0.47 \pm 2.22$  mmol/L (p = 0.57 for intervention vs control)

First author, year	Outcome incidence	Results	Covariates included in analysis
Östman E, 2006 <sup>28</sup>	Not applicable (continuous outcomes)	No changes in fasting glucose, fasting insulin and insulin sensitivity after either intervention.  No changes in insulin secretion after high glycaemic index and low fibre bread products.  Changes in insulin secretion after low glycaemic index and high fibre bread products: 1,185.4 ± 264 mIU/l min (pre-intervention) to 751.4 ± 141 mIU/l min (post-intervention) at 0-60 min (p <0.05).	None/not reported
Wein P, 1999 <sup>27</sup>	Type 2 diabetes incidence:  26.8% in intervention group and  28.1% in control group	Type 2 diabetes: relative risk for intervention vs control group: 0.63 (95% CI 0.35, 1.14).	Intervention group, entry age, entry BMI and change in BMI

First author, year	Outcome incidence	Results	Covariates included in analysis
Peterson CM, 1995 <sup>26</sup>	Not applicable (continuous outcome)	<p>Serum fasting insulin at baseline, end of first intervention, and end of second intervention, respectively (mean <math>\pm</math> standard deviation):</p> <ul style="list-style-type: none"> <li>- 40% carbohydrate group (first intervention): 17 <math>\pm</math> 6, 12 <math>\pm</math> 7, 15 <math>\pm</math> 12 mU/L</li> <li>- 55% carbohydrate group (first intervention): 24 <math>\pm</math> 13, 22 <math>\pm</math> 6, 26 <math>\pm</math> 10 mU/L</li> </ul> <p>Differences between treatment groups were not significant (<math>p &gt; 0.05</math>).</p>	None/not reported

**Table 8.** Results from observational studies included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes

First author, year	Outcome incidence	Results	Covariates included in analysis
Tobias DK, 2018 <sup>25</sup>	n = 172 diabetes cases and 175 age-matched controls (nested-case control study)	Odds ratios for type 2 diabetes for top vs bottom quartile of intake: Isoleucine: 5.78, 95% CI 1.86, 17.95 (P trend = 0.004) Leucine: 4.66, 95% CI 1.60, 13.53 (P trend = 0.01) Valine: 5.71, 95% CI 1.85, 17.60 (P trend = 0.009) Total branched-chain amino acids: 4.63, 95% CI 1.61, 13.36 (P trend = 0.01)	Age, total energy intake, fasting status $\leq$ 8 h, alcohol intake, family history of diabetes, menopausal status and current menopausal hormone therapy use, physical activity, smoking status, ethnicity, BMI, Alternative Healthy Eating Index-2010 adherence dietary quality score
Mercier R, 2018 <sup>30</sup>	n = 126 (45%) abnormal	Women with abnormal glucose tolerance had significantly lower fruit and vegetable serves (mean $\pm$ standard deviation: 6.5 $\pm$ 0.2) and	Age and BMI

First author, year	Outcome incidence	Results	Covariates included in analysis
	glucose tolerance cases, including 34% prediabetes and 11% type 2 diabetes	<p>vegetables serves (<math>3.9 \pm 0.2</math>) and tended to have lower fruit serves (<math>2.6 \pm 0.2</math>) than women with normal glucose tolerance (<math>7.4 \pm 0.2</math>, <math>4.5 \pm 0.2</math> and <math>3.0 \pm 0.1</math>, respectively) (<math>p = 0.001</math>, <math>p = 0.04</math> and <math>p = 0.10</math>, respectively).</p> <p>Odds ratios for abnormal glucose tolerance:</p> <p>Fruit and vegetables per 1 serve: 0.88, 95% CI 0.81-0.97</p> <p>Vegetables per 1 serve: 0.88, 95% CI 0.78-1.00</p> <p>Fruit per 1 serve: 0.88, 95% CI 0.76-1.02</p> <p>Fruit and vegetables <math>&lt;7</math> vs <math>\geq 7</math> serves: 1.84, 95% CI 1.13-3.00</p>	
Andersson- Hall U, 2018 <sup>31</sup>	n = 32 (23.4%) impaired glucose tolerance cases and n = 17	<p>There were no differences between groups in terms of energy, carbohydrate, protein and fat intake, meal frequency, and proportion of vegetable oil used for cooking (<math>p &gt; 0.05</math>). Compared with women with normal glucose tolerance, women with impaired glucose tolerance (<math>p = 0.01</math>) and type 2 diabetes (<math>p = 0.01</math>) had lower proportions of butter used</p>	Age, ethnicity, BMI

First author, year	Outcome incidence	Results	Covariates included in analysis
	(12.4%) type 2 diabetes cases	for cooking, and women with type 2 diabetes also had higher proportion of margarine used for cooking (p = 0.004).	
Bao W, 2016(a) <sup>22</sup>	n = 614 (16%) type 2 diabetes cases	Hazard ratios for type 2 diabetes:  Total iron intake: 1.64, 95% CI 1.20-2.25 for top (37.2 mg/d) vs bottom (11.6 mg/d) category  Supplemental iron intake: 1.83, 95% CI 1.25-2.70 for top ( $\geq 30$ mg/d) vs bottom (0 mg/d) category  Dietary haem iron intake: 1.80, 95% CI 1.18-2.74 for top (1.5 mg/d) vs bottom (0.7 mg/d) category  Dietary non-haem iron intake: 0.71, 95% CI 0.51-1.00 for top (16.6 mg/d) vs bottom (9.0 mg/d) category	Age, parity, BMI, age at first birth, ethnicity, oral contraceptive use, menopausal status, cigarette smoking, alcohol intake, physical activity, ratio of polyunsaturated fat to saturated fat, total energy intake, saturated fat, trans fat, cholesterol, animal and vegetable protein, glycaemic

First author, year	Outcome incidence	Results	Covariates included in analysis
Bao W, 2016(b) <sup>23</sup>	n = 722 (16%) type 2 diabetes cases	<p>Hazard ratios for type 2 diabetes for top vs bottom quintiles of diet scores (confounder and confounder + BMI-adjusted models, respectively):</p> <p>Overall low-carbohydrate diet score: 2.13, 95% CI 1.65-2.76 (P trend = &lt;0.001) and 1.36, 95% CI 1.04-1.78 (P trend = 0.003)</p> <p>Animal low-carbohydrate diet score: 2.18, 95% CI 1.68-2.83 (P trend = &lt;0.001) and 1.40, 95% CI 1.06-1.84 (P trend = 0.004)</p>	<p>load, cereal fibre, calcium, magnesium, vitamin C, supplemental iron (for haem and non-haem iron analysis) and total dietary iron (for supplemental iron analysis)</p> <p>Age, parity, BMI, age at first birth, ethnicity, oral contraceptive use, menopausal status, cigarette smoking, alcohol intake, physical activity, family history of diabetes, total energy intake, glycaemic index</p>

First author, year	Outcome incidence	Results	Covariates included in analysis
		Vegetable -carbohydrate diet score: 1.29, 95% CI 1.00-1.67 (P trend = 0.14) and 1.19, 95% CI 0.91-1.55 (P trend = 0.50)	
Tobias DK, 2012 <sup>24</sup>	n = 491 (11%) type 2 diabetes cases	Hazard ratios for type 2 diabetes for top vs bottom quartile of dietary pattern scores (confounder and confounder + BMI-adjusted models, respectively):  Alternate Mediterranean diet: 0.60, 95% CI 0.44-0.82 (P trend = 0.002) and 0.76, 95% CI 0.55, 1.05 (P trend = 0.13)  Dietary Approaches to Stop Hypertension: 0.54, 95% CI 0.39-0.73 (P trend = <0.001) and 0.68, 95% CI 0.49-0.94 (P trend = 0.04)  Alternate Healthy Eating Index: 0.43, 95% CI 0.31-0.59 (P trend = <0.001) and 0.65, 95% CI 0.46-0.92 (P trend = 0.01)	Age, total energy intake, parity, age at first birth, race/ethnicity, parental history of type diabetes, oral contraceptive use, menopausal status, smoking status, physical activity, alcohol intake, BMI (breastfeeding was not included in the models, but it was mentioned that this did not alter the findings)

First author, year	Outcome incidence	Results	Covariates included in analysis
Kim S-H, 2011 <sup>29</sup>	n = 161 (42.3%) prediabetes cases  n = 27 (7.1%) type 2 diabetes cases	Compared with women with normal glucose tolerance, women who developed prediabetes and type 2 diabetes had higher intakes of total energy, total and animal protein, total, animal and monounsaturated fat.  There were no differences in intake of carbohydrates, plant protein, plant, saturated, polyunsaturated and omega 3 and 6 fatty acids, and fibre.	Pre-pregnancy BMI

**[online] Table 9.** Quality assessment for each individual observational study included in a systematic review on associations of dietary intake with diabetes outcomes in women with a history of gestational diabetes<sup>a</sup>

Study	Selection		Comparability		Outcome			Total score	
	Representativeness of exposed cohort <sup>b</sup>	Selection of non-exposed cohort <sup>c</sup>	Ascertainment of exposure <sup>d</sup>	Outcome not present at start of study <sup>e</sup>	Comparability of cases and controls on bases of design or analysis <sup>f</sup>	Outcome assessment <sup>g</sup>	Adequate duration of follow-up <sup>h</sup>		Adequacy of follow-up rate <sup>i</sup>
Tobias DK, 2018 <sup>25</sup>	C	A★	B★	A★	A★ B★	B★	A★	D	7
Mercier R, 2018 <sup>30</sup>	B★	A★	B★	B	A★	A★	A★	D	6
Andersson-Hall U, 2018 <sup>31</sup>	B★	A★	B★	B	A★	A★	A★	B★	7
Bao W, 2016(a) <sup>22</sup>	C	A★	B★	A★	A★ B★	B★	A★	D	7
Bao W, 2016(b) <sup>23</sup>	C	A★	B★	A★	A★ B★	B★	A★	D	7
Tobias DK, 2012 <sup>24</sup>	C	A★	B★	A★	A★ B★	B★	A★	D	7
Kim S-H, 2011 <sup>29</sup>	A★	A★	C	A★	-	A★	A★	C	5

<sup>a</sup> Risk of bias was assessed using the Newcastle-Ottawa Scale (NOS). A study can be awarded a maximum of one star for each numbered item within the Selection and Outcome categories. A maximum of two stars can be given for Comparability. (Wells GA, Shea B, O'connell D, Peterson J, Welch V, Losos M, et al. The Newcastle-Ottawa Scale (NOS) for assessing the quality if nonrandomized studies in meta-analyses. 2009).

<sup>b</sup> Representativeness of the exposed cohort was scored as (A [star]) truly representative of the average postpartum or mid-age woman with a history of gestational diabetes in the community (random sample); (B [star]) somewhat representative of the average postpartum or mid-age woman with a history of gestational diabetes in the community; (C) selected group of users e.g. only specific occupation; (D) no description of the derivation of the cohort.

<sup>c</sup> Selection of the non-exposed cohort was scored as (A [star]) drawn from the same community as the exposed cohort; (B) drawn from a different source; (C) no description of the derivation of the non-exposed cohort.

<sup>d</sup> Ascertainment of exposure was scored as (A [star]) secure record (e.g. controlled diet); (B [star]) structured interview/validated dietary recall/diet history/ food frequency questionnaire; (C) written self-report (e.g. non-validated dietary assessment); (D) no description.

<sup>e</sup> Demonstration that outcome of interest was not present at start of study was scored as (A [star]) yes (no type 1 or type 2 diabetes prior to pregnancy and prior to study entry); (B) no.

<sup>f</sup> Comparability of cohorts on the basis of the design or analysis was scored as (A [star]) study controls for age, ethnicity and body mass index; (B [star]) study controls for any additional factor (e.g. parity, family history of diabetes, smoking, physical activity, other dietary factors).

<sup>g</sup> Outcome assessment was scored as (A [star]) independent blind assessment (e.g. hospital record/oral glucose tolerance test); (B [star]) record linkage or validated self-reported diagnosis; (C) self-report (not validated); (D) no description.

<sup>h</sup> Adequate duration of follow-up for outcomes to occur was scored as (A [star]) yes, if – after baseline – all women were screened for type 2 diabetes or answered a question on whether they were diagnosed or treated for type 2 diabetes; (B) no.

<sup>i</sup> Adequacy of follow-up rate was scored as (A [star]) complete follow up or all subjects accounted for using multiple imputation; (B [star]) subjects lost to follow up unlikely to introduce bias (samples are comparable based on comparison of in- and excluded participants) or  $\leq 10\%$  lost during follow-up; (C) follow up rate  $< 90\%$  or no description of those lost; (D) no statement.