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# A randomized feasibility trial of time-restricted eating during pregnancy in people with increased risk of gestational diabetes

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Time-restricted eating (TRE) is a nutritional intervention that confines the daily time-window for energy intake. TRE reduces fasting glucose concentrations in non-pregnant individuals, but whether this eating protocol is feasible and effective for glycemic control in pregnancy is unknown. The aim of this randomized controlled trial was to investigate the adherence to and effect of a 5-week TRE intervention (maximum 10 h daily eating window) among pregnant individuals at risk of gestational diabetes mellitus (GDM), compared with a usual-care control group. Participants underwent 2-h oral glucose tolerance tests and estimation of body composition, before and after the intervention. Interstitial glucose levels were continuously measured, and adherence rates and ratings of hunger were recorded daily. Thirty of 32 participants completed the trial. Participants allocated to TRE reduced their daily eating window from 12.3 (SD 1.3) to 9.9 (SD 1.0) h, but TRE did not affect glycemic measures, blood pressure, or body composition, compared with the control group. TRE increased hunger levels in the evening, but not in the morning, and induced only small changes in dietary intake. Adhering to a 5-week TRE intervention was feasible for pregnant individuals with increased risk of GDM but had no effect on cardiometabolic outcomes.

Keywords Diet, Continuous glucose monitoring, Insulin, Obesity, Female, Prevention

The prevalence of diabetes is increasing in parallel with the obesity pandemic, with gestational diabetes mellitus (GDM) estimated to occur in up to 14% of all pregnancies<sup>1</sup>. GDM is the development of glucose intolerance with onset or first recognition during pregnancy, brought on by an underlying chronic insulin resistance due to beta-cell dysfunction<sup>2,3</sup>. Important risk factors for GDM include advanced maternal age, a body mass index (BMI)  $\geq$  25 kg/m<sup>2</sup>, previous GDM or a family history with diabetes, delivery of a macrosomic child, and non-white ethnicity<sup>4,5</sup>. Glucose intolerance during pregnancy increases the risk of adverse pregnancy outcomes, such as pre-eclampsia, pre-term birth, macrosomia, caesarean delivery, and birth injury<sup>2,6,7</sup>. Furthermore, GDM increases the risk of developing diabetes and cardiovascular disease later in life for both the mother and the offspring<sup>2,7,8</sup>.

Lifestyle interventions, including nutritional therapy, is regarded as the primary strategy for managing GDM<sup>5,9</sup>. However, there is no consensus on which diet is best for achieving optimal glycemic control in pregnancy<sup>5,10</sup>. Time-restricted eating (TRE) is a dietary strategy in which the time-window for energy intake each day is restricted, typically to between 6 and10 h/day. TRE has shown to have positive effects on glucose regulation in people with overweight/obesity<sup>11,12</sup> and to improve body composition<sup>13</sup>. As such, TRE has emerged as a potentially beneficial intervention for individuals at risk of, or diagnosed with GDM<sup>14</sup>. However, we are unaware of published research on the feasibility or effect of TRE in pregnant individuals. The primary aim of this randomized controlled trial (RCT) was to examine the adherence to TRE in pregnant individuals with at least one risk factor for developing GDM. Secondary outcomes included the effect of TRE on markers of metabolic

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### Results Participants

Thirty-two participants undertook baseline assessments and were randomized to either TRE (n = 15) or a control group (CON, n = 17) (Fig. 1). The first participant was recruited 18/01/2019 and the last date of follow-up was 13/03/2023, with a halt in inclusion of participants from March to September 2020 during the Covid pandemic. We stopped the inclusion when we had reached our pre-specified number of participants. Two individuals withdrew their consent to participate in the study after baseline testing and randomization. All participants had at least one risk factor for GDM according to the Norwegian recommendations for screening of GDM<sup>15</sup>, with some having more than one risk factor. Most of the included participants (n = 27) had a pre-pregnancy BMI  $\geq 25 \text{ kg/m}^2$ . Nineteen were expecting their first child at an age  $\geq 25 \text{ years}$ , two were of Asian or African ethnicity, three had first-degree relatives with diabetes, two had previously given birth to a child with birthweight > 4.5 kg, and one had been diagnosed with GDM in a previous pregnancy. Table 1 shows an overview of baseline characteristics according to group.

#### Time-restricted eating was feasible in pregnancy and affected feelings of hunger

The participants allocated to TRE reduced their daily eating window from 12.0 h (SD 2.2) at baseline to 9.9 h (SD 1.6) during the 5-week intervention period, with no change in eating window in CON in the same period (12.5 h (SD 1.9) at baseline and 13.1 h (SD 1.7) in the following 5 weeks) (Fig. 2). During the baseline week, the participants in the TRE group consumed their first meal at 09:17 h (SD 2.0) and their last meal at 21:01 h (SD 1.1), whereas the average times for the first and last meal during the intervention were 08:43 h (SD 1.9) and 18:50 h (SD 1.7), respectively (Fig. 3). Individuals in the TRE group adhered to the  $\leq$  10-h eating window on 4.7 (SD 0.4) days/week during the intervention period, giving an adherence rate of 67% (SD 6%).



Fig. 1. Flow diagram of participants in the study.

	n	Control $(n = 17)$	n	Time-restricted eating $(n = 15)$
Age, years	17	30.1 (2.9)	1 (2.9) 15 32.2 (3.8)	
Gestational week	17	19.8 (5.2)	15	18.6 (6.7)
Primipara/Multipara, number	17	11/6	15 8/7	
Height, cm	17	169 (7) 15 163 (9)		163 (9)
Weight, kg	17	81.3 (15.2)	15	79.6 (16.4)
Body mass index, kg/m <sup>2</sup>	17	28.5 (5.3)	15	29.9 (5.5)
Muscle mass, kg	17	28.5 (4.1)	15	26.9 (4.2)
Fat mass, kg	17	29.7 (11.0)	15	30.8 (11.7)
Fat percentage	17	35.6 (7.5)	15	37.8 (7.3)
Visceral fat area, cm <sup>2</sup>	17	140 (54)	15	147 (60)
Systolic BP, mmHg	16	115 (7)	15	113 (10)
Diastolic BP, mmHg	16	76 (5)	15	74 (6)
Heart rate, beats/min	16	77 (12)	15	80 (10)
Fasting glucose, mmol/L	17	4.5 (0.4)	15	4.5 (0.3)
120-min glucose, mmol/L	17	4.9 (1.3)	15	5.3 (1.8)
Fasting insulin, µlU/mL	17	14.2 (5.4)	15	16.5 (8.7)
120-min insulin, µlU/mL	17	42.0 (29.0)	15	69.8 (97.4)
HbA1c, mmol/mol	17	29.8 (3.1)	14	31.0 (3.4)
HOMA2-IR	17	1.7 (0.6)	15	2.0 (1.0)
Cholesterol, mmol/L	17	5.4 (1.0)	13	5.7 (0.9)
Triglycerides, mmol/L	17	1.5 (0.4)	13	1.5 (0.5)
HDL, mmol/L	17	2.0 (0.4)	13	2.0 (0.4)
LDL, mmol/L	17	3.4 (0.8)	13	3.7 (0.8)

**Table 1**. Baseline characteristics of participants, according to group allocation. Data are means with standarddeviation (SD) if not otherwise stated. BP = Blood pressure, HDL = High-density lipoprotein cholesterol,HOMA2-IR = Homeostatic model assessment of insulin resistance, LDL = Low-density lipoprotein cholesterol.



**Fig. 2.** Mean eating window duration. Observed values at baseline and during the 5-week intervention period according to group. Descriptive statistics with standard deviation for the intention-to-treat population. p – value was computed using linear mixed model, comparing the time-restricted eating (TRE) group with the control group (CON).

There were no effects of the TRE intervention on the participants' ratings of hunger in the morning (Supplementary Fig. S1, Supplementary Table S1). In the evening during the second week of the intervention, the participants in TRE reported increased hunger levels, desire to eat, and prospective intake of energy, and decreased satiety, compared with CON (Supplementary Fig. S2, Supplementary Table S1). Evening hunger was



**Fig. 3**. Individual changes in eating window for participants allocated to time-restricted eating. Lines show the average self-reported window for energy intake per day for each participant in the intervention group. The graph is based on observed values from participant handbooks.

still higher and evening satiety lower in the TRE group in the last week of the intervention, compared with CON, but the desire to eat in the evening was not.

#### Time-restricted eating had no effect on secondary metabolic outcomes

At baseline, none of the participants fulfilled the Norwegian criteria for GDM: fasting glucose between 5.3 and 6.9 mmol/L and/or 120-min glucose between 9.0 and 11.0 mmol/L after a 75-g oral glucose tolerance test (OGTT)<sup>15</sup>. Testing post-intervention revealed that one participant in CON had fasting glucose of 5.7 mmol/L, with all remaining participants being below the threshold for GDM. TRE had no effect on any of the glycemic or metabolic outcomes (Table 2), nor on 24-h glucose area under the curve (AUC), day-time glucose AUC or night-time glucose AUC (Fig. 4, Supplementary Table S2). The TRE intervention had no effect on systolic or diastolic blood pressure (Table 2).

# Time-restricted eating had small effect on dietary intake and no effect on physical activity

TRE had no effect on total energy intake (Supplementary Table S3). Participants in the TRE group consumed 33 g less carbohydrates per day in the last week of the intervention period, compared with CON (p=.040). In the second week of the intervention period, participants in the TRE group consumed 34 g less sugar compared with CON (p=.005), but there was no difference between groups in the last week of the intervention. There were no between-group differences in other dietary intake variables (Supplementary Table S3), or in estimated total energy expenditure, or other measures of physical activity throughout the study (Supplementary Table S4).

#### Adverse events

No adverse events were reported during the study.

		Baseline After the intervention		Difference (group x time) compared with CON				
Outcome	Group	n	Mean (SD)	n	Mean (SD)	Est. effect	95% CI	p
Fasting glucose, mmol/L	CON	17	4.5 (0.4)	16	4.5 (0.4)			
	TRE	15	4.5 (0.3)	14	4.4 (0.2)	-0.1	-0.3 to 0.1	0.187
120-min glucose, mmol/L	CON	17	4.9 (1.3)	16	5.0 (1.6)			
	TRE	15	5.3 (1.8)	14	5.6 (1.6)	0.2	-0.4 to 0.8	0.529
HbA1c, mmol/mol	CON	17	29.8 (3.1)	16	30.4 (4.2)			
	TRE	14	31.0 (3.4)	14	31.2 (3.9)	-0.3	-2.3 to 1.3	0.811
Fasting insulin, µlU/ml	CON	17	14.2 (5.4)	16	15.8 (6.6)			
	TRE	15	16.5 (8.7)	14	16.9 (9.3)	-0.7	-4.6 to 3.3	0.742
120-min insulin, μlU/ml	CON	17	42.0 (29.0)	16	53.7 (42.0)			
	TRE	15	69.8 (97.4)	13	90.3 (72.6)	12.1	-29.3 to 43.0	0.670
HOMA2-IR	CON	17	1.7 (0.6)	16	1.9 (0.8)			
	TRE	15	2.0 (1.0)	14	2.1 (1.1)	-0.1	-0.5 to 0.4	0.743
Muscle mass, kg	CON	17	28.5 (4.1)	16	29.2 (4.7)			
	TRE	15	26.9 (4.2)	14	27.2 (4.2)	-0.5	-1.0 to 0.1	0.091
Fat mass, kg	CON	17	29.7 (11.0)	16	28.9 (8.4)			
	TRE	15	30.8 (11.7)	14	32.6 (11.6)	0.2	-0.7 to 1.2	0.609
Weight, kg	CON	17	81.3 (15.2)	16	81.6 (13.7)			
	TRE	15	79.6 (16.4)	14	82.0 (16.3)	-0.4	-1.3 to 0.5	0.383
Visceral fat area, cm <sup>2</sup>	CON	17	139.5 (53.7)	16	138.2 (46.1)			
	TRE	15	146.7 (60.2)	14	153.6 (59.4)	-1.1	-7.3 to 5.0	0.707
Systolic BP, mmHg	CON	16	114.9 (6.7)	15	113.5 (5.5)			
	TRE	15	113.0 (9.7)	14	109.7 (6.0)	-3.0	-7.0 to 1.0	0.142
Diastolic BP, mmHg	CON	16	75.5 (5.3)	15	74.4 (4.7)			
	TRE	15	74.0 (5.8)	14	71.4 (6.0)	-2.4	-5.4 to 0.6	0.117
Resting heart rate, bpm	CON	16	77.3 (12.3)	15	77.4 (10.8)			
	TRE	15	80.1 (10.2)	14	79.0 (8.5)	-0.1	-5.3 to 5.1	0.982
Total cholesterol, mmol/L	CON	17	5.4 (0.9)	14	6.0 (1.0)			
	TRE	13	5.7 (0.9)	12	6.3 (0.9)	0.0	-0.3 to 0.3	0.856
HDL-cholesterol, mmol/L	CON	17	2.0 (0.4)	14	2.1 (0.4)			
	TRE	13	2.0 (0.4)	13	2.1 (0.4)	0.0	-0.2 to 0.2	0.783
LDL-cholesterol, mmol/L	CON	17	3.4 (0.8)	14	3.9 (0.9)			
	TRE	13	3.7 (0.8)	13	4.4 (0.9)	0.0	-0.3 to 0.3	0.936
Triglycerides, mmol/L	CON	17	1.5 (0.4)	14	1.8 (0.5)			
	TRE	13	1.5 (0.5)	13	2.1 (0.7)	0.1	-0.1 to 0.3	0.394

**Table 2.** Intention-to-treat analyses of secondary outcomes. Baseline data are reported as mean values with standard deviations (SD) of observed values at baseline, and post-intervention for *n* participants in each group. Results from linear mixed model analysis presents estimated effect, which represents the difference in mean in the time-restricted eating group (TRE) compared with the control group (CON), with corresponding 95% confidence interval (CI) and *p*-values. BP=blood pressure, BPM=beats per minute, HbA1c=haemoglobin A1c, HDL=high-density lipoprotein, HOMA2-IR=homeostatic assessment of insulin resistance, LDL=low-density lipoprotein.

#### Discussion

This study was the first experimental investigation of adherence rates to a TRE intervention during pregnancy. Our results largely support our main hypothesis that 5 weeks of TRE is feasible during pregnancy, as the participants adhered to a  $\leq$  10 h/day eating window on ~5 days/week throughout the intervention period. The average eating window was just under 10 h/day in the intervention period, representing a 2-h reduction from baseline. Despite this reduced eating window, we failed to detect any beneficial effects of TRE on glycemic control or other metabolic outcomes.

The adherence rate to TRE in our study was 67%, which is somewhat lower than the rates reported in other studies of TRE involving non-pregnant individuals and with a daily eating window of maximum 10  $h^{13,16}$ . We have previously showed that reproductive-aged women with overweight/obesity managed to adhere to an identical TRE intervention on 6.2 days/week (89%) for 7 weeks<sup>13</sup>, while Anton and colleagues reported an adherence rate of 84% to a similar 4-week intervention among overweight, sedentary adults aged 65 years or older<sup>16</sup>. All these studies had a relatively short intervention period, but Lin et al.<sup>17</sup> reported 87% adherence to an 8-h TRE protocol over the course of a 12-month study in participants with obesity. The main reason for



**Fig. 4**. Area under curve (AUC) glucose. (a) 24-hours AUC, (b) day-time AUC, and (c) night-time AUC. Data are estimated from continuous glucose monitoring using Glyculator 3.0. Symbols show averages and error bars show standard deviations for the control group (CON) and the time-restricted eating group (TRE) in the baseline week (Week 1), the first three weeks of the intervention period (Weeks 2–4), and in the last two weeks of the intervention period (Weeks 5–6). *p* – values are for between-group comparisons using linear mixed models.

lower adherence in the present study is likely the pregnant state of our participants. In pregnancy, nausea and preference of specific foods (cravings) are common, especially in the first trimester. These factors may affect the ability to consume energy only at specific periods during the day as required for TRE. Indeed, a qualitative study on attitudes towards TRE among people who were pregnant or had recently given birth, reported that some were concerned about the baby's health, nausea, and hunger<sup>14</sup>. However, 47% of the participants in that study perceived TRE as safe during pregnancy, but only 24% of them said they would be willing to try a TRE regimen during pregnancy<sup>14</sup>.

We report no serious adverse effects of adhering to a 10-h TRE window for 5 weeks and neither do previous studies<sup>11,13,16-20</sup>. The most commonly reported adverse effects of TRE are nausea, headaches, dizziness, diarrhea, and dry mouth. However, these effects will either diminish over time, or are resolved by increasing water intake<sup>11,16,18</sup>. Despite that a 10-h TRE window was found to be feasible in the current study, including participants earlier in pregnancy could yield different results. The mean gestational age in the intervention group in our study was 18.6 weeks (ranging from 12 to 30 weeks) at baseline, whereas nausea is most common in early pregnancy<sup>21</sup>. The on-going BEFORE THE BEGINNING trial will determine the feasibility and effectiveness of TRE also in early pregnancy<sup>22</sup>.

There was no effect of TRE on any of the measured cardiometabolic outcomes in our study. A reason for this could be that 5 weeks may be insufficient to induce any significant glycemic changes in pregnant individuals<sup>23</sup>. However, in non-pregnant individuals, TRE interventions do improve glycemic control after interventions of similar duration<sup>13,18,24</sup>. In one study, restricting the window for energy intake to between 08:00 and 16:00 h improved skeletal muscle insulin sensitivity among healthy males after 2 weeks<sup>24</sup>. Similar results were also seen in a crossover trial involving men with prediabetes, in which 5 weeks of TRE with a 6-h eating window early in the day reduced the concentrations of insulin both in the fasting state and during a 3-h OGTT<sup>18</sup>. In both these studies, the window of energy intake was substantially shorter than in the present study, which may explain the different findings. In our previous study involving reproductive-aged women with overweight/obesity, we showed lower nocturnal glucose after 7 weeks of TRE with a 10-h window for energy intake <sup>13</sup>. In contrast to our previous trial, the participants in the present study did not reduce their energy intake during the intervention period.

Concurrent with no change in energy intake or expenditure, 5 weeks of TRE did not affect body weight. Weight loss is often the goal of TRE interventions<sup>11,25-27</sup>, with improvements in glycemic measures frequently occurring after weight loss<sup>13</sup>. Even if most TRE regimens allow unrestricted intake of energy within the stipulated eating window, people typically reduce their daily energy intake unintentionally, which likely underpins most of the cardiometabolic benefits of TRE<sup>28</sup>. The weight loss observed after TRE interventions makes it difficult to determine whether there is a weight loss-independent effect of TRE. However, in the study by Sutton and colleagues, in which the participants were provided with standardized meals to maintain energy balance and weight, they observed several improvements in glycemic outcomes<sup>18</sup>. Conversely, others have reported reductions in body weight after a TRE intervention without concomitant improvements in measures of glycemic control<sup>27</sup>. Additionally, we found no change in blood pressure or resting heart rate after the intervention, which is in concordance with findings in similar trials<sup>11,13,27</sup>. Conversely, some trials have reported reduced blood pressure following TRE interventions<sup>18,29-31</sup>.

We placed no restrictions on the timing of the eating window, but we recommended that the participants started their daily energy intake by 09:00 h. On average, participants commenced their daily eating window at 08:43 h and ended it at 18:50 h during the intervention. Placing the eating window early in the day has been shown to improve glycemic outcomes in animals and humans<sup>18,32,33</sup>, and it is suggested to be beneficial for glucose metabolism and insulin sensitivity to synchronize meal timing with the circadian rhythm. Implementing an eating schedule synchronized with the body's natural activity-rest cycles has been shown to lower blood glucose and insulin concentrations<sup>33,34</sup>. In pregnant individuals, it was recently shown that consuming > 50% of total daily energy intake between 19:00 and 07:00 h was associated with less desirable glycemic outcomes,

including increased fasting glucose and higher 24-h glucose levels<sup>35</sup>. A previous study by the same research group also indicated that increased maternal night-fasting intervals were associated with decreased fasting glucose in the late-second trimester of pregnancy<sup>36</sup>. Furthermore, in non-pregnant individuals with overweight or obesity, early TRE has superior effects compared with later TRE in improving glycemic control<sup>37</sup>.

The participants in our study had increased risk of developing GDM, but none had signs of abnormal glucose metabolism at baseline. A more pronounced effect of TRE on glycemic outcomes would be more likely if the participants' glucose metabolism was compromised<sup>23</sup>. As such, TRE has demonstrated improved cardiometabolic outcomes in people with impaired glucose metabolism, including individuals with the metabolic syndrome, pre-diabetes, or type 2 diabetes<sup>18,19,31,38,39</sup>.

There are several limitations to this study and the interpretation of results. We had relatively few participants and the intervention period was only 5 weeks. We did not adjust the p-values for multiple comparisons and, therefore, there is a risk of type 1 error. However, several of the between-group changes we observed in ratings of hunger were highly significant, with p-values < 0.001. The participants who volunteered for our study were probably less bothered by nausea than the general population of pregnant people with increased risk of GDM. We recruited participants in either the second or third semester of pregnancy (mean gestational week 19, range 12–30 weeks). While it would have been ideal to recruit all participants at the same stage of pregnancy to investigate the effect of TRE on glycemic outcomes and body weight changes, this was not practical given the constraints of the study. However, our main aim was to assess the adherence to TRE during pregnancy as a measure of its feasibility among pregnant individuals. Due to the randomized design of the study, there was no systematic difference between the groups in gestational length or any other baseline characteristics at the time of inclusion. Furthermore, even if all the participants in our study had increased risk of GDM, not all had a BMI above the healthy range. Since the physiological responses to a dietary intervention may differ according to baseline BMI, future studies should investigate whether TRE can limit gestational weight gain in people with elevated BMI.

It is also possible that the intervention period was too short, that duration of the daily eating window was too long, or that the eating window was placed too late in the day, to impact glycemic control and other cardiometabolic outcomes. Considering the unique physiological state of pregnancy, our results are difficult to compare with studies in non-pregnant populations. Pregnant individuals require special considerations and adjustments, and the metabolic response in pregnant people could differ significantly from that of non-pregnant individuals.

#### Conclusion

To the best of our knowledge, the present study is the first trial to determine the adherence to TRE in pregnancy. While the results of our study suggest that TRE is feasible in the second and third trimesters of pregnancy among individuals at risk of GDM, there were no improvements in glycemic or cardiometabolic outcomes in our subject cohort. Further studies should include a larger number of participants and implement TRE beyond 5 weeks to determine the long-term feasibility and potential health benefits of TRE during pregnancy.

#### Methods

#### Study design and participants

This parallel-group RCT was carried out at the Norwegian University of Science and Technology (NTNU) and St. Olav's hospital. The study was approved by the Regional Committees for Medical and Health Research Ethics in Norway (ID 12366) and registered in Clinical Trials (ClinicalTrials.gov, identifier: NCT03803072, 14/01/2019). The study was performed in accordance with the Declaration of Helsinki. All participants provided written informed consent before participating in the study. Participants were recruited through public advertising at St. Olav's hospital, on social media, and university web pages. We screened for eligibility via telephone before we included participants in the trial. Table 3 shows the inclusion and exclusion criteria. The study period for each participant was 6 weeks, including one week of baseline measurements and 5 weeks of TRE for the intervention group or no intervention for participants in CON. Fasting venous blood sampling, OGTT, estimation of body composition, blood pressure and heart rate measurements were undertaken at the baseline visit and after the 5-week intervention period (Fig. 5). After baseline assessments were completed, the participants were randomly allocated (1:1) to either 5 weeks of TRE or usual care (CON). All participants were fitted with continuous glucose monitors (CGMs) and physical activity monitors and received instructions on how to self-report their daily eating window in the study handbook. Dietary intake and physical activity were measured for 7 days in the baseline week, in the second week of the intervention, and in the last week of the intervention (Fig. 5).

Inclusion criteria	Exclusion criteria
<ul> <li>Age ≥ 18 years</li> <li>Pregnant with a singleton foetus in gestational week 12–30</li> <li>Understands written and spoken Norwegian or English</li> <li>At least one risk factor for gestational diabetes (GDM) as defined by Norwegian guidelines for GDM screening<sup>15</sup>:</li> <li>o Pre-pregnancy body mass index ≥ 25 kg/m<sup>2</sup></li> <li>o First birth at the age of ≥ 25 years</li> <li>o Previous GDM</li> <li>o Previous delivery of newborn ≥ 4.5 kg</li> <li>o First degree relative with diabetes mellitus</li> <li>o Asian or African ethnicity</li> </ul>	<ul> <li>Habitual daily eating window ≤ 12 h</li> <li>Previously diagnosed with diabetes type 1 or 2</li> <li>Shift work that includes nightshifts</li> </ul>

#### Table 3. Inclusion and exclusion criteria.



**Fig. 5**. Study design. Participants visited the laboratory for assessments at baseline before randomization, and after the intervention. These visits included fasting blood sampling, an oral glucose tolerance test, estimation of body composition, blood pressure and heart rate measurements. The participants wore continuous glucose monitors throughout the study and physical activity monitors in the baseline week, week 3, and week 6. They registered the time points for first and last energy intake each day throughout the study and reported dietary intake in the baseline week, week 3, and week 6. baseline week of the study.

## **Randomization and blinding**

The participants undertook baseline assessments before being randomly allocated (1:1) to either TRE or CON, after stratification for BMI < 27 kg/m<sup>2</sup> or  $\geq$  27 kg/m<sup>2</sup>. The first or the last author performed the randomization using a random number generator (WebCRF, The Unit for Applied Clinical Research, NTNU, Trondheim). The randomization sequence was concealed in the WebCRF until interventions were assigned. Neither participants nor study personnel were blinded.

#### Intervention

All participants continued with their habitual eating habits during the baseline week. The participants allocated to TRE were instructed to limit their time-window for energy intake to maximum 10 h/day for 5 weeks. We advised the participants to start their eating window no later than 9:00 h and to consume their last energy intake no later than 19:00 h. They could freely consume energy-free beverages such as black coffee, tea, and diet soda outside their daily eating window. We gave no advice regarding the amount of energy or types of foods to be consumed. Participants allocated to CON were instructed to continue their habitual eating pattern for the entire 6-week study period. All participants received a booklet about healthy lifestyle habits in pregnancy.

#### Primary outcomes

The primary outcome measure in this study was the adherence to TRE during pregnancy. Adherence to was self-reported in a study handbook, in which the participants recorded the time points of their first and last energy intake every day throughout the whole study period. We calculated adherence as the average duration of the daily eating window throughout the 5-week study period, as well as the average number of days per week the participants adhered to the  $\leq$  10-h eating window.

#### Secondary outcomes

#### Blood analyses

We sampled venous blood at baseline and after 6 weeks. The participants attended the laboratory after a 10-h overnight fast. After the fasting blood sample was obtained, the participants underwent a 120-min OGTT in which they ingested 75 g glucose dissolved in 250 mL water (GlucosePro, Norges Naturmedisinsentral AS). A second venous blood sample was obtained 120 min after the ingestion of the glucose solution. Glycemic outcomes include fasting plasma glucose, 120-min plasma glucose after the OGTT, HbA1c, fasting serum insulin,

and 120-min serum insulin after the OGTT. Additionally, fasting blood total cholesterol, triglycerides, highdensity lipoprotein (HDL) and low-density lipoprotein (LDL) were measured. EDTA tubes were centrifuged immediately after sampling at 2220G and 4 °C for 10 min. Serum tubes rested upright for 30 min before being centrifuged at 2220G and 20 °C for 10 min. All analyses apart from insulin concentrations were carried out at the laboratory at St. Olav's hospital. Additional aliquots of plasma, serum and full blood were stored at -80 C for later analysis. We analyzed fasting and 120-min serum insulin using enzyme-linked immunosorbent assay (ELISA, IBL-International, Hamburg, Germany). These analyses were carried out per manufacturer's instructions using a DS2 ELISA processing system (Dynex Technologies, Virginia, USA) at the Department of Circulation and Medical Imaging, NTNU. Using fasting blood glucose levels and fasting insulin levels, we calculated insulin resistance (HOMA2-IR) using the online HOMA2 calculator (https://www.rdm.ox.ac.uk/about/our-clinicalfacilities-and-mrc-units/DTU/software/homa).

#### Continuous glucose monitoring

The participants wore CGMs (FreeStyle Libre 1, Abbott Diabetes Care, Norway) throughout the entire 6-week study period. They were fitted with CGM sensors at baseline and instructed to do minimum four scans evenly spaced out throughout each day of the study. We gave out replacement sensors at baseline to cover the whole 6-weeks study duration. We covered the screen of the CGM monitor to avoid that the participants got aware of their glucose levels. Raw CGM data were processed using Microsoft Excel and divided into a 1-week baseline period (week 1), a 3-week mid-study period (weeks 2–4) and 2-week end-period (weeks 5–6). We used the Glyculator 3.0 calculator (https://glyculator.btm.umed.pl) to impute missing glucose measurements due to infrequent scans and to estimate glucose AUC for 24 h, daytime (06:00–00:00) and night-time (00:01–05:59).

#### Body composition, blood pressure, and resting heart rate

We estimated the participants' body composition using a bioelectrical impedance scale (InBody770, Biospace CO, Ltd, Seoul, Korea) after  $a \ge 10$  h overnight fast. The participants wore light clothing and no shoes during these tests. Parameters used to estimate body composition include height, weight, BMI, fat mass, muscle mass and visceral fat area. Blood pressure and heart rate were measured using an automated blood pressure device (Welch Allyn, Germany). We obtained three measurements with 1-min intervals and report the average of these three measurements.

#### Physical activity and diet

We fitted the participants with physical activity monitors (BodyMedia Sensewear Armband, Pittsburgh, PA) during the baseline testing. These monitors were worn during the baseline week, the 3rd study week, and the 6th study week to estimate average weekly physical activity and energy expenditure. Raw physical activity data were processed using Microsoft Excel and divided into baseline (week 1), mid-study (weeks 2–4), and end-period (weeks 5–6). In the same periods, the participants recorded their dietary intake for 7 days using an online food diary (kostholdsplanleggeren.no). Raw dietary data were processed using Microsoft Excel. In the same weeks, participants recorded subjective feelings of hunger, fullness, satiety, and desire to eat in the mornings and evenings on 10-cm visual analogue scales printed in their study handbooks.

#### Sample size

We did not perform a formal power analysis for this trial. Generally, a sample size between 24 and 50 is recommended to estimate effect size and standard deviation in feasibility studies<sup>40-42</sup>. We aimed to include 32 participants to ensure that we had a minimum of 24 (12 in each group) with measurements of glucose tolerance and insulin sensitivity at two time points, accounting for an expected drop-out of 20%.

#### Statistical analysis

The adherence data are reported as descriptive statistics. We additionally used linear mixed models (LMM) to compare adherence in TRE with CON. In the LMM models, we included time and the interaction between time and group as fixed factors, and participants as random factor<sup>43</sup>. We report the estimated mean change in the TRE group, with corresponding 95% confidence intervals (CI) and p-values, compared with CON. We also used linear mixed models to compare secondary outcome measures between TRE and CON. The normality of residuals was checked by visual inspection of QQ-plots. For variables that were not normally distributed (HbA1c, insulin 120 min, HDL), we performed bootstrapping with 3000 samples (bias corrected and accelerated CIs). All randomized participants were included in the intention-to-treat analysis, regardless of adherence. We excluded data with less than 4 days of valid CGM measurements or physical activity data in a period, as well as less than at least two weekdays and one weekend day of nutritional intake. However, if the participant had sufficient data in other time periods, then these data were included in the analyses. Statistical analyses were performed in IBM SPSS Statistics 27. We consider p-values <0.05 as statistically significant and have not performed any adjustments for multiple comparisons due to the exploratory nature of our research questions.

#### Data availability

Data reported in this paper will be shared by the corresponding author upon reasonable request.

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#### References

- Wang, H. et al. Estimation of global and regional gestational diabetes mellitus prevalence for 2021 by international association of diabetes in pregnancy study group's criteria. *Diabetes Res. Clin. Pract.* 183, 109050. https://doi.org/10.1016/j.diabres.2021.109050 (2022).
- Buchanan, T. A., Xiang, A. H. & Page, K. A. Gestational diabetes mellitus: risks and management during and after pregnancy. *Nat. Rev. Endocrinol.* 8, 639–649. https://doi.org/10.1038/nrendo.2012.96 (2012).
- Plows, J. F., Stanley, J. L., Baker, P. N., Reynolds, C. M. & Vickers, M. H. The pathophysiology of gestational diabetes mellitus. Int. J. Mol. Sci. https://doi.org/10.3390/ijms19113342 (2018).
- Zhang, C., Rawal, S. & Chong, Y. S. Risk factors for gestational diabetes: is prevention possible? *Diabetologia*. 59, 1385–1390. https://doi.org/10.1007/s00125-016-3979-3 (2016).
- ACOG Practice Bulletin No. 190: gestational diabetes Mellitus. Obstet. Gynecol.131, e49-e64. https://doi.org/10.1097/ aog.00000000002501 (2018).
- Johns, E. C., Denison, F. C., Norman, J. E. & Reynolds, R. M. Gestational diabetes Mellitus: mechanisms, treatment, and complications. *Trends Endocrinol. Metab*.29, 743–754. https://doi.org/10.1016/j.tem.2018.09.004 (2018).
- Sullivan, S. D., Umans, J. G. & Ratner, R. Gestational diabetes: implications for cardiovascular health. *Curr. Diab Rep.* 12, 43–52. https://doi.org/10.1007/s11892-011-0238-3 (2012).
- Szmuilowicz, E. D., Josefson, J. L. & Metzger, B. E. Gestational diabetes mellitus. *Endocrinol. Metab. Clin. North. Am.*48, 479–493. https://doi.org/10.1016/j.ecl.2019.05.001 (2019).
- Moholdt, T., Hayman, M., Shorakae, S., Brown, W. J. & Harrison, C. L. The role of lifestyle intervention in the prevention and treatment of gestational diabetes. *Semin Reprod. Med.* https://doi.org/10.1055/s-0040-1722208 (2021).
- 10. Yamamoto, J. M. et al. Gestational diabetes Mellitus and Diet: a systematic review and Meta-analysis of randomized controlled trials examining the impact of modified dietary interventions on maternal glucose control and neonatal Birth Weight. *Diabetes Care.* **41**, 1346–1361. https://doi.org/10.2337/dc18-0102 (2018).
- 11. Cienfuegos, S. et al. Effects of 4- and 6-h time-restricted feeding on weight and cardiometabolic health: a randomized controlled trial in adults with obesity. *Cell. Metab*. **32**, 366-378e363. https://doi.org/10.1016/j.cmet.2020.06.018 (2020).
- 12. Parr, E. B., Devlin, B. L., Radford, B. E. & Hawley, J. A. A delayed morning and earlier evening time-restricted feeding protocol for improving glycemic control and dietary adherence in men with overweight/obesity: a randomized controlled trial. *Nutrients* (2020).
- Haganes, K. L. et al. Time-restricted eating and exercise training improve HbA1c and body composition in women with overweight/ obesity: a randomized controlled trial. *Cell. Metab.* 34, 1457–1471e1454. https://doi.org/10.1016/j.cmet.2022.09.003 (2022).
- Flanagan, E. W., Kebbe, M., Sparks, J. R. & Redman, L. M. Assessment of eating behaviors and perceptions of time-restricted eating during pregnancy. J. Nutr. 152, 475–483. https://doi.org/10.1093/jn/nxab397 (2022).
- Helsedirektoratet Svangerskapsdiabetes. Nasjonal faglig retningslinje for svangerskapsdiabetes, (2018). https://helsedirektoratet.no/ retningslinjer/svangerskapsdiabetes
- Anton, S. D. et al. The effects of Time restricted feeding on overweight, older adults: a pilot study. Nutrients. 11https://doi. org/10.3390/nu11071500 (2019).
- Lin, S. et al. Time-restricted eating without calorie counting for weight loss in a racially diverse population: a randomized controlled trial. Ann. Intern. Med. 176, 885–895. https://doi.org/10.7326/m23-0052 (2023).
- 18. Sutton, E. F. et al. Early time-restricted feeding improves insulin sensitivity, blood pressure, and oxidative stress even without weight loss in men with prediabetes. *Cell. Metab*.27, 1212-1221e1213. https://doi.org/10.1016/j.cmet.2018.04.010 (2018).
- Pavlou, V. et al. Effect of time-restricted eating on weight loss in adults with type 2 diabetes: a Randomized Clinical Trial. JAMA Netw. Open.6, e2339337. https://doi.org/10.1001/jamanetworkopen.2023.39337 (2023).
- Suthutvoravut, U. et al. Efficacy of time-restricted eating and behavioral economic intervention in reducing fasting plasma glucose, hba1c, and cardiometabolic risk factors in patients with impaired fasting glucose: a randomized controlled trial. *Nutrientshttps://* doi.org/10.3390/nu15194233 (2023).
- 21. Festin, M. Nausea and vomiting in early pregnancy. BMJ Clin. Evid. 2014 (2014).
- Sujan, M. A. J. et al. Randomised controlled trial of preconception lifestyle intervention on maternal and offspring health in people with increased risk of gestational diabetes: study protocol for the before the beginning trial. *BMJ Open.*13, e073572. https://doi. org/10.1136/bmjopen-2023-073572 (2023).
- Kang, J. et al. Effect of time-restricted feeding on Anthropometric, metabolic, and Fitness parameters: a systematic review. J. Am. Nutr. Assoc. 41, 810–825. https://doi.org/10.1080/07315724.2021.1958719 (2022).
- Jones, R. et al. Two weeks of early time-restricted feeding (eTRF) improves skeletal muscle insulin and anabolic sensitivity in healthy men. Am. J. Clin. Nutr. 112, 1015–1028. https://doi.org/10.1093/ajcn/ngaa192 (2020).
- 25. Moro, T. et al. Effects of eight weeks of time-restricted feeding (16/8) on basal metabolism, maximal strength, body composition, inflammation, and cardiovascular risk factors in resistance-trained males. *J. Transl Med.* **14**, 290. https://doi.org/10.1186/s12967-016-1044-0 (2016).
- Jamshed, H. et al. Early time-restricted feeding improves 24-hour glucose levels and affects markers of the circadian clock, aging, and autophagy in humans. *Nutrientshttps://doi.org/10.3390/nu11061234* (2019).
- Chow, L. S. et al. Time-restricted eating effects on body composition and metabolic measures in humans who are overweight: a feasibility study. Obes. (Silver Spring)28, 860–869. https://doi.org/10.1002/oby.22756 (2020).
- Chang, Y., Du, T., Zhuang, X. & Ma, G. Time-restricted eating improves health because of energy deficit and circadian rhythm: a systematic review and meta-analysis. *iScience*. 27, 109000. https://doi.org/10.1016/j.isci.2024.109000 (2024).
- Gabel, K. et al. Effects of 8-hour time restricted feeding on body weight and metabolic disease risk factors in obese adults: a pilot study. Nutr. Healthy Aging. 4, 345–353. https://doi.org/10.3233/nha-170036 (2018).
- Zhang, L. M. et al. Randomized controlled trial for time-restricted eating in overweight and obese young adults. *iScience*. 25, 104870. https://doi.org/10.1016/j.isci.2022.104870 (2022).
- Wilkinson, M. J. et al. Ten-hour time-restricted eating reduces weight, blood pressure, and atherogenic lipids in patients with metabolic syndrome. *Cell. Metab.* 31, 92–104e105. https://doi.org/10.1016/j.cmet.2019.11.004 (2020).
- 32. Jamshed, H. et al. Early time-restricted feeding improves 24-hour glucose levels and affects markers of the circadian clock, aging, and autophagy in humans. *Nutrients*11, (2019).
- Chaix, A., Lin, T., Le, H. D., Chang, M. W. & Panda, S. Time-restricted feeding prevents obesity and metabolic syndrome in mice lacking a circadian clock. *Cell. Metab.*29, 303–319e304. https://doi.org/10.1016/j.cmet.2018.08.004 (2019).
- Chaix, A., Manoogian, E. N. C., Melkani, G. C. & Panda, S. Time-restricted eating to prevent and manage chronic metabolic diseases. *Annu. Rev. Nutr.*39, 291–315. https://doi.org/10.1146/annurev-nutr-082018-124320 (2019).
- Loy, S. L. et al. Associations of predominant night-eating with plasma glycemic status and continuous glucose monitoring measures among pregnant women. *Clin. Nutr.*42, 2320–2327. https://doi.org/10.1016/j.clnu.2023.10.009 (2023).
- Loy, S. L. et al. Maternal circadian eating time and frequency are associated with blood glucose concentrations during pregnancy. J. Nutr.147, 70–77. https://doi.org/10.3945/jn.116.239392 (2017).
- Liu, J., Yi, P. & Liu, F. The effect of early time-restricted eating vs later time-restricted eating on weight loss and metabolic health. J. Clin. Endocrinol. Metab. 108, 1824–1834. https://doi.org/10.1210/clinem/dgad036 (2023).
- Che, T. et al. Time-restricted feeding improves blood glucose and insulin sensitivity in overweight patients with type 2 diabetes: a randomised controlled trial. Nutr. Metab. (Lond). 18, 88. https://doi.org/10.1186/s12986-021-00613-9 (2021).

- 39. Hutchison, A. T. et al. Time-restricted feeding improves glucose tolerance in men at risk for type 2 diabetes: a randomized crossover trial. *Obes. (Silver Spring).* 27, 724–732. https://doi.org/10.1002/oby.22449 (2019).
- Lancaster, G. A., Dodd, S. & Williamson, P. R. Design and analysis of pilot studies: recommendations for good practice. J. Eval Clin. Pract. 10, 307–312. https://doi.org/10.1111/j.2002.384.doc.x (2004).
- 41. Browne, R. H. On the use of a pilot sample for sample size determination. Stat. Med.14, 1933-1940 (1995).
- 42. Sim, J. & Lewis, M. The size of a pilot study for a clinical trial should be calculated in relation to considerations of precision and efficiency. *J. Clin. Epidemiol.***65**, 301–308. https://doi.org/10.1016/j.jclinepi.2011.07.011 (2012).
- J, T. et al. Different ways to estimate treatment effects in randomised controlled trials. Contemp. Clin. Trials Commun. 10, 80–85. https://doi.org/10.1016/j.conctc.2018.03.008 (2018).

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# Author contributions

T.M., K.Å.S., and J.A.H. conceived the study and analysis plan. H.M.S.S., K.L.H., T.M.G., and M.K.J. collected the data. H.M.S.S., M.A.J.S., K.L.H., and T.M. analyzed the data. All authors contributed to the interpretation of the data. H.M.S.S. drafted the manuscript. All authors critically reviewed the manuscript for intellectual content and gave their final approval of the version to be published.

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# Declarations

# **Competing interests**

The authors declare no competing interests.

# Additional information

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