A single serving of caffeinated coffee impairs postprandial glucose metabolism in overweight men

Tracey M. Robertson1, Michael N. Clifford1, Simon Penson2, Gemma Chope2 and M. Denise Robertson1*

1Department of Nutritional Sciences, University of Surrey, Guildford GU2 7WG, UK
2Department of Primary Production and Processing, Campden BRI, Chipping Campden GL55 6LD, UK

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Abstract

Previous studies regarding the acute effects of coffee on glycaemic control have used a single large dose of coffee, typically containing the caffeine equivalent of 2–4 servings of coffee. This study investigates whether the acute effects of coffee are dose-dependent, starting with a single serving. A total of ten healthy overweight males participated in a two-part randomised double-blind cross-over study. In the first part, they ingested 2, 4 or 8 g instant decaffeinated coffee (DC) dissolved in 400 ml water with caffeine added in proportion to the DC (total 100, 200 or 400 mg caffeine) or control (400 ml water) all with 50 g glucose. In the second part, they ingested the same amounts of DC (2, 4, 8 g) or control, but with a standard 100 mg caffeine added to each. Capillary blood samples were taken every 15 min for 2 h after each drink and glucose and insulin levels were measured. Repeated measures ANOVA on glucose results found an effect when caffeine was varied in line with DC (P=0.008). Post hoc analysis revealed that both 2 and 4 g DC with varied caffeine content increased the glycaemic response v. control. There was no effect of escalating doses of DC when caffeine remained constant at 100 mg. These results demonstrate that one standard serving of coffee (2 g) is sufficient to affect glucose metabolism. Furthermore, the amount of caffeine found in one serving (100 mg) is sufficient to mask any potential beneficial effects of increasing other components. No dose-dependent effect was found.

Key words: Human: Coffee: Caffeine: Glucose: Insulin: Metabolism: Postprandial metabolism

According to the WHO, nearly 350 million people worldwide have diabetes, of whom approximately 90 % have type 2 diabetes mellitus (T2DM)(31). If poorly controlled, diabetes can result in serious complications such as CVD, nephropathy and retinopathy. Much epidemiological evidence suggests that people who drink coffee have a reduced risk of developing T2DM(2–5). This has been demonstrated for both caffeinated and decaffeinated coffee (DC)(6,7) and appears to be particularly strong for those who drink large quantities of coffee(8,9), however it is not known precisely how coffee may exert this protective effect. Metabolic syndrome is a group of risk factors that are linked with the development of T2DM. Potential mechanisms where coffee drinking could reduce the risk of T2DM may be through its demonstrated effects on some of these factors. These include aiding weight loss by increasing energy expenditure and fat oxidation(10–12) and reducing energy intake(13), increasing HDL-cholesterol(14) and increasing circulating adiponectin levels(15,16). Adiponectin levels have been shown to be inversely associated with insulin resistance(17), therefore, an increase in levels may result in improved insulin sensitivity, although this has not been found previously, perhaps due to short study durations.

Acute studies about the effects of coffee or caffeine on the postprandial glycaemic response have, however, generally shown a detrimental effect. Greater glucose and insulin AUC have been demonstrated following caffeine ingestion(18,19). Caffeinated coffee ingestion has also produced increased glucose responses compared with a control(20–22); however, the effect appears to be less pronounced than that produced by caffeine alone(23). It has been suggested that bioactive components in coffee, other than caffeine, may be attenuating the documented acute negative effects of caffeine on glycaemia. Chlorogenic acids (CGA) and quinides have attracted attention as possible candidates with suggested mechanisms including delayed glucose absorption by the enterocyte(24) via dissipation of the Na+ gradient and consequent inhibition of the SGLT1 glucose transporter(25) in addition to increased uptake of glucose in non-skeletal muscles(26). Studies involving CGA on its own(27) and coffee enriched with CGA(28) have both demonstrated a reduction in glucose response v. control. The varying results from coffee studies may be partially explained by the

Abbreviations: CGA, chlorogenic acids; CQA, caffeoylquinic acid; DC, decaffeinated coffee; iAUC, incremental AUC; T2DM, type 2 diabetes mellitus.

* Corresponding author: Dr M. D. Robertson, fax +44 1483 688501, email m.robertson@surrey.ac.uk
CGA content of the coffees used, as this can vary considerably\(^{29}\). However, this cannot be verified as the majority of these studies did not measure CGA content. A limitation to previously published work is the relatively high dose of caffeine used, typically 3–6 mg caffeine/kg body weight\(^{19,30}\), which equates to 210–420 mg for a 70 kg person. A Canadian study\(^{33}\) that analysed the caffeine content of various instant coffees, prepared to participants' personal taste preferences, found a median caffeine concentration in the prepared coffees of 328 µg/ml (range 102–559 µg/ml). This median value would provide 85 mg caffeine in a standard-sized (260 ml) serving. Doses of 3–6 mg/kg would, therefore, potentially be equivalent to 2.5–5 standard-sized servings of instant coffee taken as a single dose for a 70 kg person and correspondingly higher for heavier individuals.

To our knowledge, there has been no study thus far investigating the glycaemic effects of a single serving of instant coffee as consumed in the UK. Instant coffee was chosen for this study because (i) it is the most commonly consumed type of coffee in the UK\(^{32}\) and (ii) we wanted to ensure that the levels of the different coffee components were consistent between brews as other types of coffee have been shown to vary in their caffeine and CGA content\(^{34,35}\). In this pilot investigation, we aimed to perform a dose–response study in overweight individuals with the lowest dose set at a single serving of instant coffee as consumed in the UK. In Part B of the study, CGA and other coffee components were increased step-wise, whereas the caffeine dose remained constant, to investigate the acute effects of fortification with these other coffee components.

**Methods**

**Participants**

Ten participants were recruited from the staff and student population of the University of Surrey. All of them were overweight (BMI > 25 kg/m\(^2\)) but otherwise healthy adult males aged 19–62 years who were regular coffee drinkers (>4 servings/week). All of them were non-smokers. Participants were required to have been weight stable for the previous 3 months and to have no history of heart disease, diabetes, liver disease or any gastrointestinal (GI) or endocrine disorders. This study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects were given a favourable ethical opinion by the University of Surrey Ethics Committee. Written informed consent was obtained from all the participants. The study was carried out between August 2012 and February 2013.

**Study design**

A randomised cross-over design was used to investigate the glycaemic effect of three different doses of DC, to which different doses of caffeine had been added. A control drink (water). All drinks were made up in 400 ml of water and each had 50 g glucose powder added. The different drink combinations along with their caffeoylequinic acid (CQA) contents are summarised in Table 1. All treatments, apart from control, were double-blinded. Participants were informed that they were testing different coffee drinks only; they were not told that they contained differing amounts of caffeine and coffee, DC, to which caffeine powder was added, was used so that the effects of caffeine and the other coffee components could be examined separately. The instant coffee granules were made and centrifuged at 1500 g for 10 min, and the plasma was transferred into serum microvette tubes. Plasma glucose concentrations were measured immediately using the YSI 2300.

**Test protocol**

All the participants received each of the eight test beverages in a random order on separate study mornings separated by a washout period of at least 5 d. An online randomisation programme was used to generate the sequences for each participant. For 2 d before each of the study days, the participants were asked to refrain from exercise, alcohol and all coffee- and caffeine-containing foods and drinks. They were asked to maintain the same diet the day before each visit and were given a standardised meal to consume the evening before. The standard meal was a supermarket macaroni cheese ready meal. All measurements were taken after a 12-h fast. On each study day, the participants arrived in the morning and provided a fasted capillary blood sample via the finger-prick technique. They then consumed the test drink within 10 min and provided further capillary blood samples every 15 min for 2 h. They were asked to rate each drink using a scale of 1–5 for four drink quality measures: strength, palatability, taste and smell.

**Biochemistry**

All blood samples were collected by the finger-prick method using a lancing device and collected into glucose (heparin fluoride) microvette tubes (Sarstedt)\(^{36}\). Approximately 300 µl of blood was collected at each time point. These were refrigerated and centrifuged at 1500 g for 10 min, and the plasma was transferred into serum microvette tubes. Plasma glucose concentrations were measured immediately using the YSI 2300.
STAT Plus™ (YSI Life Sciences). Inter-assay CV were 5% and intra-assay CV were 2% for the glucose analysis. Following glucose analysis, the plasma was frozen at –20°C until the end of the study for batch analysis. Plasma insulin concentrations were measured by colorimetric sandwich ELISA (EMD Millipore). Inter-assay CV were 7% and intra-assay CV were 3% for the insulin analysis. The finger-prick technique was used as the blood collected via this method is primarily arterial blood and consequently reflects the absorbed glucose levels more accurately than venous blood. Venous blood glucose values are usually lower than arterial values in the postprandial state(35) as some of the absorbed blood glucose will have been taken up by skeletal muscles before it gets to the sample site.

Calculations and statistical analysis
Statistical analysis was carried out using IBM SPSS Statistics 19 (IBM). Sample size was set at ten participants, in line with standard practise in glycaemic index testing(36). The incremental AUC (iAUC) for glucose and insulin for each treatment was calculated by the trapezoid method, with the area under baseline being ignored as recommended by the FAO(37) and as described by Brouns et al.(36). The Matsuda Index (10,000/√(fasting glucose × fasting insulin) × (mean glucose × mean insulin during OGTT))38 was calculated as a measure of postprandial insulin sensitivity. Overall treatment effects were analysed by repeated measures ANOVA. A Bonferroni adjustment for multiple comparisons was applied when post hoc analysis was used. Statistical significance was taken as P<0.05. All results are expressed as mean values with their standard error of the mean, apart from participant characteristics and drink questionnaire results, which are expressed as mean values and standard deviations.

Results
All the participants were overweight males with a mean age of 30·4 (SD 14·2) years, BMI of 27·8 (SD 2·2) kg/m², fasting blood glucose levels of 5·2 (SD 0·5) mmol/l and were all insulin sensitive (fasting range 5·2–29 pmol/l)(39). There was no significant difference between visits for fasting glucose and insulin measures. All were regular coffee drinkers with a mean caffeinated coffee intake of 2·3 (SD 1·6) servings/d.

Drink quality
Mean responses for each of the four drink quality measures – strength, palatability, taste and smell – are shown in Table 2. Repeated measures ANOVA tests were performed for Parts A and B. For Part A, an overall significant difference was found for smell (P=0·045) with no significant differences between individual pairs. No other differences were found in Part A. For Part B, an overall significant difference was found for palatability

| Table 1. Composition of the eight coffee drinks used in this study* |
|------------------|------------------|------------------|------------------|------------------|------------------|
| Drink number | Decaffeinated granules (g) | Added caffeine (mg) | Total caffeine (mg) | Total CQA (mg) | Glucose (g) | Water (ml) |
| Part A | 1 | 0 | 0 | 0 | 0 | 50 | 400 |
| 2 | 2 | 94 | 100 | 47 | 50 | 400 |
| 3 | 4 | 188 | 200 | 94 | 50 | 400 |
| 4 | 8 | 376 | 400 | 188 | 50 | 400 |
| Part B | 5 | 0 | 100 | 0 | 50 | 400 |
| 6 | 2 | 94 | 100 | 47 | 50 | 400 |
| 7 | 4 | 88 | 100 | 94 | 50 | 400 |
| 8 | 8 | 76 | 100 | 188 | 50 | 400 |

CQA, caffeoylquinic acid.
* Decaffeinated coffee and caffeine powder were dissolved in 200 ml boiling water with 50 g glucose powder added; 200 ml of cold water was then added to allow quick consumption.

Fig. 1. Diagrammatic representation of the relative proportions of decaffeinated coffee (DC) to caffeine (CF) for the eight test drinks used in this study. In Part A, CF rises in proportion to DC, whereas in Part B CF remains constant at 100 mg. --- instant DC; •••••• CF.
(P=0.020) and taste (P=0.036) with no significant differences between individual pairs. A significant difference in strength (P<0.001) was detected in Part B, with the post hoc analysis revealing drink 8 to have been perceived as stronger than both drinks 6 (P=0.040) and 5 (P=0.002). A difference in smell (P=0.025) was also found in Part B, with the post hoc analysis revealing a difference between drinks 5 and 6 (P=0.001).

Glucose and insulin

For Part A, where the amount of caffeine was varied in line with the DC dose, there was a significant difference between treatments for glucose. Repeated measures ANOVA on the glucose time points showed a significant treatment effect (P=0.008) and a time × treatment effect (P=0.022). Post hoc analysis showed a significant difference between the control drink (drink 1 in Table 1) and both the one- and two-serving equivalents (drinks 2 and 3) (P<0.032). A significant difference was also found for both iAUC (P=0.019) and peak values (P=0.006), with the post hoc analysis showing a significant difference between drinks 1 and 2 for both iAUC (P=0.008) and peak values (P=0.006) and a trend for a difference between drinks 1 and 3 for peak value (P=0.056). The mean peak blood glucose concentration for the control (drink 1) was 8.25 mmol/l, whereas the peak for Part A was 9.62 mmol/l (for the two servings equivalent: drink 3), a mean increase of 17% (1.37 mmol/l) in peak values. No other differences between treatments were detected. There was no difference between treatments for postprandial insulin or insulin sensitivity as measured by the Matsuda Index. The 2-h postprandial glucose and insulin responses for Part A are shown in Fig. 2. The mean iAUC and peak values for glucose and insulin for Parts A and B are shown in Table 3.

For Part B of the study, where all drinks contained 100 mg caffeine, a significant difference in glucose was found for peak values only (P=0.049), with no difference between any

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### Table 2. Drink questionnaire results*

<table>
<thead>
<tr>
<th>Drink number</th>
<th>Palatability</th>
<th>Strength</th>
<th>Taste</th>
<th>Smell</th>
</tr>
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<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Part A</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>1</td>
<td>2.9</td>
<td>1.3</td>
<td>2.6</td>
<td>1.1</td>
</tr>
<tr>
<td>2</td>
<td>3.9</td>
<td>0.6</td>
<td>2.7</td>
<td>0.7</td>
</tr>
<tr>
<td>3</td>
<td>3.7</td>
<td>0.8</td>
<td>3.3</td>
<td>0.7</td>
</tr>
<tr>
<td>4</td>
<td>3.0</td>
<td>1.2</td>
<td>3.3</td>
<td>0.9</td>
</tr>
<tr>
<td>Part B</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>2.7</td>
<td>1.1</td>
<td>2.1a</td>
<td>1.1</td>
</tr>
<tr>
<td>6</td>
<td>3.7</td>
<td>0.7</td>
<td>2.5a</td>
<td>0.8</td>
</tr>
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<td>7</td>
<td>4.1</td>
<td>0.6</td>
<td>3.1</td>
<td>0.7</td>
</tr>
<tr>
<td>8</td>
<td>3.8</td>
<td>0.8</td>
<td>3.6b</td>
<td>0.8</td>
</tr>
</tbody>
</table>

* Mean values within a column/part with unlike superscript letters were significantly different (P<0.05).

* Participants rated the control and coffee drinks on a scale of 1–5 for palatability, strength, taste and smell.
individual pairs in the post hoc analysis. There were no significant differences for insulin responses or insulin sensitivity between treatments, although there was a time \times treatment effect for insulin ($P = 0.013$). The 2-h postprandial glucose and insulin responses for Part B are shown in Fig. 3.

A paired $t$ test comparing the glucose iAUC for the two control drinks (1 and 5) showed a significant difference between the two ($P = 0.045$), with the control drink containing caffeine (drink 5) having a 22% higher mean iAUC. There were no significant differences between the two control drinks for any other measures.

### Discussion

Our study adds to the body of evidence from previous acute studies demonstrating a temporary worsening in glucose response following coffee ingestion. We have shown, for the first time, that a single serving of instant caffeinated coffee, as typically consumed in the UK, is sufficient to disrupt the 2-h postprandial glucose response. Furthermore, Part B of our study demonstrates that the amount of caffeine found in one serving of coffee can attenuate any possible beneficial effects of escalating doses of the other coffee components.

Gavieli et al.\(^{40}\) used two different amounts of caffeinated coffee, 3 and 6 mg/kg body weight (mean: 228 and 457 mg), which were similar to the two higher amounts of caffeine used in Part A (200 and 400 mg, drinks 3 and 4) and found similar results. In line with our study, they reported a higher glucose iAUC in both coffee doses compared with control, with no significant dose–response effect. Although they did not find any overall intervention effect for the glucose response, they did find a time \times treatment effect. Our post hoc analysis found the increased glucose response only in the two lower doses of coffee plus caffeine (drinks 2 and 3) and not in the highest dose. It is unlikely that a greater insulin response to the high caffeine levels had suppressed the glucose response in this case, as we found no significant differences between drinks for the insulin response. Furthermore, although Beaudoin et al.\(^{41}\) did find a dose–response effect on insulin, they did not find a direct suppression of glucose as a result. We also observed a difference between treatments in the timing of the peak glucose values, with

### Table 3. Mean incremental AUC (iAUC) and peak glucose and insulin values for the 2-h postprandial period (Mean values and standard deviations)

<table>
<thead>
<tr>
<th>Drink number</th>
<th>Decaffeinated granules (g)/total caffeine (mg)</th>
<th>Glucose iAUC Mean</th>
<th>Glucose iAUC SD</th>
<th>Peak glucose (mmol/l) Mean</th>
<th>Peak glucose (mmol/l) SD</th>
<th>Insulin iAUC Mean</th>
<th>Insulin iAUC SD</th>
<th>Peak insulin (pmol/l) Mean</th>
<th>Peak insulin (pmol/l) SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Part A</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>0/0</td>
<td>172(^{a})</td>
<td>89</td>
<td>8.9(^{a})</td>
<td>1.3</td>
<td>8238</td>
<td>6114</td>
<td>193</td>
<td>109</td>
</tr>
<tr>
<td>2</td>
<td>2/100</td>
<td>237(^{e})</td>
<td>102</td>
<td>10.0(^{b})</td>
<td>1.4</td>
<td>10,294</td>
<td>7376</td>
<td>239</td>
<td>139</td>
</tr>
<tr>
<td>3</td>
<td>4/200</td>
<td>243</td>
<td>108</td>
<td>9.6</td>
<td>1.3</td>
<td>8521</td>
<td>5141</td>
<td>206</td>
<td>111</td>
</tr>
<tr>
<td>4</td>
<td>8/400</td>
<td>218</td>
<td>117</td>
<td>9.5</td>
<td>1.4</td>
<td>10,098</td>
<td>7848</td>
<td>224</td>
<td>97</td>
</tr>
<tr>
<td>Part B</td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>0/100</td>
<td>210(^{c})</td>
<td>77</td>
<td>8.9</td>
<td>1.3</td>
<td>6094</td>
<td>4374</td>
<td>151</td>
<td>74</td>
</tr>
<tr>
<td>6</td>
<td>2/100</td>
<td>237</td>
<td>104</td>
<td>9.8</td>
<td>1.5</td>
<td>6887</td>
<td>4523</td>
<td>159</td>
<td>61</td>
</tr>
<tr>
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<td>110</td>
<td>9.6</td>
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<td>1.1</td>
<td>7584</td>
<td>5845</td>
<td>205</td>
<td>84</td>
</tr>
</tbody>
</table>

\(^{a,b,c}\) Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).
the control drink peaking at 15 min while all the others peaked later at 30 min. This is consistent with the theory that coffee may exert a beneficial effect by delaying glucose absorption.

In Part B of our study, we found no overall treatment effect even at the highest dose of DC (8 g), suggesting that even a relatively small dose of caffeine (100 mg, equivalent to that found in one serving of caffeinated coffee) may be enough to override any potential beneficial effects of increasing other coffee compounds. We did see a difference in peak glucose values across the four treatments in Part B; however, the study was not powered to detect pair-wise difference in individual treatments. The control drink in Part B also contained 100 mg caffeine, unlike in Part A where it was simply water plus glucose, which would explain why there was no significant difference between the control drink and any of the coffee drinks. When the two control drinks were compared, we found the iAUC to be significantly higher for the caffeine-containing control drink (5), suggesting that it is indeed caffeine that is causing the increased glycaemic response in Part A. It should be noted that the CGA content of coffees can vary considerably, depending on bean composition, processing and brewing method, and a detailed analysis of our coffee samples found it to have a relatively low total CQA content of 23.5 mg/g (CQA being the largest proportion of CGA in coffee). This is equivalent to approximately 47 mg in our lowest dose coffee and 188 mg in our highest dose (Table 1). This is low compared with espresso-type coffees, which have a mean CQA content of 145 mg/serving (range 24–422 mg)

Part A, however, is comparable to the mean CQA content of six brands of instant DC, previously measured at 24–4 mg/g.

Nevertheless, these relatively low levels in our coffee may partially explain why we saw no effect of escalating doses of CGA in Part B of our study.

We did not demonstrate any dose–response effect of coffee on blood glucose or insulin levels, either when the caffeine was increased along with the coffee (Part A) or when the caffeine was kept constant (Part B). This is in contrast with a recent study that found a dose–response effect when using caffeine alone; however, the caffeine-only study had more participants (n = 24), and – as noted previously – caffeine has been shown to produce a greater effect than coffee on post-prandial glucose and insulin levels. The lack of a dose–response effect may also be due to the high inter- and intra-individual variations in glycaemic response, which was apparent when individuals’ responses were examined and is in line with that previously demonstrated. Our relatively low number of participants (ten), although being the recommended minimum number for GI testing, may not be sufficient to determine a dose–response relationship. A post hoc power calculation revealed our study to be underpowered to detect a difference in insulin levels. In order for the detected differences in insulin to be statistically significant at 80% power, we would have needed more than twenty-three participants. This is based on the result that found 5/10 subjects had an increase in insulin levels after coffee intake compared with control and 5/10 subjects had a decrease in insulin levels, and thus no consistent pattern in response was observed. The doses given to our participants were not adjusted according to their body weight in contrast with the previously discussed studies as we wanted to reflect normal nutritional intake patterns. When converted to milligram caffeine per kilogram body weight, our lowest dose (100 mg caffeine) gave a mean value of 1.2 mg/kg (range 0.9–1.5 mg/kg) and our highest dose (400 mg caffeine) was 4.6 mg/kg (range 3.7–6.1 mg/kg). This variation in dose between participants may be a confounder in determining a dose–response relationship; however, it does not detract from the main result, which is that a single cup of coffee as drunk in the UK increases the post-prandial glycaemic response. Another possible confounder may be different genetic polymorphisms in the CYP1A2 gene that have been shown to affect the rate of caffeine metabolism in humans.

Unfortunately, our participants were not genotyped for these polymorphisms as this may have partly explained the high inter-individual variations we found. An obvious difference between the control and coffee drinks is the presence/absence of coffee with its characteristic bitter taste. The hedonic properties of food/drink have been shown to impact acutely on postprandial nutrient handling, and thus it was important to exclude this as a simple pre-ingestive mechanism. Interestingly, the only drink quality difference from Part A (where a postprandial glucose difference was found) was with ‘smell’; there was no overall impact on palatability. This would indicate that the impact of coffee on glucose metabolism was due to a true post-ingestive mechanism. In light of our results, and those of others discussed previously, it would be advisable for establishments carrying out GI testing not to offer coffee as an accompaniment to the test food.

In conclusion, we have demonstrated for the first time that a single serving of coffee significantly increases the postprandial glycaemic response in healthy overweight men. What is not known is whether this statistically significant increase in blood glucose is physiologically relevant and the implications for individuals with abnormal glucose tolerance. Although we observed a high percentage difference in peak values, it was quickly resolved in insulin-sensitive individuals with all treatments, including control, displaying similar responses from 60 min onwards. It is important that further investigations are carried out in those with impaired fasting glycaemia and T2DM where the effects of coffee may be clinically relevant. It should be noted that this was an acute study and its results cannot be translated simply into chronic effects. Longer-term interventions investigating these chronic effects are now needed if we are to confirm the beneficial effects of coffee suggested by the epidemiology.

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