Almond "Appetizer" Effect on Glucose Tolerance Test (GTT) Results

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Background: The extent to which glucose intolerance can be acutely improved with dietary modification is unclear. The purpose of this study was to test the effect of ingesting a low-calorie almond preload ("appetizer") 30 minutes before oral glucose tolerance testing in glucose-intolerant individuals without diabetes.

Methods: Twenty adults with prediabetes or isolated 1-hour glucose ≥ 160 mg/dL underwent 2 fasting oral glucose tolerance tests (GTTs)—1 standard GTT and 1 GTT 30 minutes after eating a half ounce (12) of dry-roasted almonds. Fourteen participants met 1 or more prediabetes diagnostic criteria; 6 had only elevated 1-hour glucose ≥ 160 mg/dL.

Results: The mean 1-hour plasma glucose after the almond preload was 37.1 mg/dL (19.4%) lower (154.6 vs 191.7; P < .001) than in the standard GTT. The almond preload reduced the area under the glucose curve by 15.5% (P < .001). Eight individuals had a marked hypoglycemic effect (glucose reduced by 45 to 110 mg/dL); 4 had a moderate hypoglycemic effect (22–32 mg/dL).

Conclusion: A low-calorie almond "appetizer" showed promise as an option for decreasing postprandial hyperglycemia in individuals with prediabetes or isolated 1-hour postprandial hyperglycemia. Further study is needed to confirm and refine the role of such a premeal appetizer in the self-care of prediabetes. (J Am Board Fam Med 2016;29:759–766.)

Keywords: Area Under Curve, Blood Glucose, Eating, Energy Intake, Fasting, Food Habits, Glucose Intolerance, Glucose Tolerance Test, Hyperglycemia, Prediabetic State, Prunus dulcis, Self-Care

Nearly 30 million Americans (almost 10% of adults) had diabetes in 2012 (mostly type 2), and another 86 million Americans (almost 30% of adults) had prediabetes.¹ Patients known to have prediabetes are at high risk for developing diabetes.² Although exercise, weight control, and hypoglycemic medications have been shown to prevent progression from prediabetes to diabetes,³ many patients are unwilling or unable to make good use of these options.

Progression from normal glucose tolerance to diabetes is associated with a progressive decrease in

acute insulin response to ingested carbohydrates and a variable reduction in insulin sensitivity. Both dysfunction and loss of pancreatic β -cells contribute to glucose intolerance.⁴ Toxic effects of hyperglycemia on β -cells contribute to the progression from impaired glucose tolerance to diabetes.⁵

One of the main causes of postprandial hyperglycemia is reduced or delayed insulin release in response to nutrient ingestion.⁶ During an oral glucose tolerance test (GTT), first-phase insulin response begins within 5 minutes after glucose ingestion and lasts for about 30 minutes. With prediabetes, some individuals have reduced or absent first-phase insulin response, whereas others have normal first-phase insulin concentrations but do not maintain normal glucose concentrations because of reduced insulin sensitivity. The second phase of acute insulin response begins 10 to 15 minutes after glucose ingestion and persists while the hyperglycemic stimulus is present.⁶

During a GTT, glucose and insulin concentrations typically peak at 30 to 45 minutes. Glucose

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intolerance at 1 hour on the GTT has serious clinical implications, even when the 2-hour glucose is <140 mg/dL. Elevated 1-hour glucose concentrations are the earliest, and often the only, reflection of impaired first-phase insulin release. Glucose intolerance at 1 hour seems to be the earliest physiologic indicator of an individual's proneness to type 2 diabetes.⁶ One-hour GTT >155 mg/dL predicts future development of diabetes better than abnormal fasting or 2-hour GTT values.⁷

Lifestyle approaches to reducing hyperglycemia have focused mainly on improving insulin sensitivity (via exercise and weight loss) or limiting total carbohydrate intake. The extent to which glucose intolerance can be improved with dietary modification is unclear, but several studies have shown that manipulating the timing of part of one's caloric intake can improve glucose tolerance.

Heacock et al⁸ found that ingesting 10 g of fructose 30 or 60 minutes before eating a highcarbohydrate test meal reduced the positive incremental area under the glucose curve by 25% to 27%. A soya-yogurt snack (30 g of soybeans and 75 g of yogurt) 2 hours before breakfast reduced the postprandial glucose increment by almost 40%.⁹ Josse et al¹⁰ and Jenkins et al¹¹ showed a 21% to 42% reduction in peak postprandial hyperglycemia by adding 30 to 90 g of almonds to a test meal containing 50 g of carbohydrates. Tan and Mattes¹² found that 43 g (1.5 oz) of almonds given as midmorning and midafternoon snacks mildly lowered postprandial serum glucose.

The purpose of this study was to test the effect of a low-calorie dietary preload (12 dry-roasted almonds, eaten 30 minutes in advance of a GTT) on GTT results in individuals with prediabetes and/or an elevated glucose concentration on a 1-hour GTT. The study hypothesis was that "priming the pancreatic pump" with a low-calorie "appetizer" 30 minutes before GTT would significantly decrease the hyperglycemic response at 1 hour in glucose-intolerant individuals who do not have diabetes. In theory, it would do so by boosting insulin secretion before ingesting the main caloric load. A prestudy analysis of glucose-intolerant patients' GTTs at the study site found that the mean 1-hour GTT value was 40 mg/dL higher than the 2-hour GTT value (185 vs 145 mg/dL), showing more room for improvement in the 1-hour than the 2-hour value.

The one-half ounce (14.2 g) almond snack given in this study contained 84 kcal, including 3 kcal from 0.7 g carbohydrate, 12 kcal from 4 g protein, and 68 kcal from 7.5 g fat (mostly polyunsaturated, averaging 65% omega-6 linoleic acid and 27% omega-3 α -linolenic acid).¹³ If the study results support the hypothesis, glucose-intolerant individuals who respond well to the almond preload and regularly ingest such a premeal appetizer might (1) have lower peak glucose concentrations after eating, and (2) be hyperglycemic less of the time.

Compared with published studies of the effects of almonds on glucose tolerance, this study entailed eating a smaller amount of almonds (½ vs 1–3 oz) and with different timings: 30 minutes before, rather than 2 hours before or during, main calorie consumption. Prefeeding the low-calorie (84 kcal) almond serving with "appetizer" timing may produce a beneficial effect on glucose intolerance similar to that seen in studies of prefeeding fructose (44 kcal)⁹ or soya-yogurt (180 kcal).¹⁰

Methods

In this crossover experimental study, each participant served as their own control. Inclusion criteria were age 18 to 90 years, prediabetic level of HbA_{1c} (5.7 to 6.4%) and elevated (\geq 160 mg/dL) 1-hour glucose on a standard fasting 2-hour GTT. Exclusion criteria included diabetes diagnosis (fasting GTT result >125 mg/dL and/or 2-hour result >200 mg/dL); currently taking any medication affecting glucose concentration (including insulin, other hypoglycemic medications, and oral or injected corticosteroids); history of almond allergy; inability to render informed consent because of cognitive or other impairment; inability or unwillingness to complete 2 oral GTTs within 4 weeks; and severe acute or unstable chronic illness.

Study participants, comprising a convenience sample, were initially identified by searching the study site's electronic medical record (Centricity) using the above-mentioned inclusion criteria. Potentially eligible patients were contacted via E-mail or regular mail, then by telephone, to elicit their interest in participating and to screen for the above-mentioned exclusion criteria. Those who still seemed to be eligible and expressed interest were scheduled for an initial study visit, when written informed consent was obtained. Those whose 1-hour result on a standard GTT was >160 mg/dL were invited to complete a second GTT (after almond ingestion).

Each full study participant underwent 2 fasting oral GTTs, most of them 2 to 7 days apart. One was a standard 75-g GTT, and the other was a GTT that began 30 minutes after ingesting a half ounce of dry-roasted almonds (12 average-sized almonds) and drinking an 8-ounce glass of tap water. Each almond load was weighed as exactly half an ounce on a food scale, and it was ingested in a mean time of 3.2 minutes. Venous blood was sampled for serial measurements of plasma glucose at baseline, and at 1 and 2 hours after each 75-g glucose load. Initially the order of the standard GTT and the almond GTT was varied randomly to control for the sequence effect. However, since a large percentage of initial enrollees did not meet the inclusion criterion of 1-hour glucose $\geq 160 \text{ mg/}$ dL, the standard GTT was done first in 17 of the 20 participants.

To minimize the variability of GTT results, participants were asked to eat as they usually did throughout the study. Participants were asked to refrain from vigorous physical activity for 8 hours before each GTT and to minimize their physical activity during each GTT. Participants completed the 14-item Perceived Stress Scale¹⁴ and a 24-hour food, alcohol, and activity record the day before each GTT.

The study was conducted in a university-affiliated, community-based family medicine residency program clinic. Blood was drawn by 1 of several phlebotomists in the on-site laboratory and sent to an outside laboratory for analysis. Blood samples from 16 of the participants were analyzed by the affiliated hospital laboratory; blood samples from 4 participants were analyzed by 1 commercial laboratory that the patients' managed Medicare insurance required.

A sample size calculation used the standard deviation for the 1-hour glucose (33 mg/dL) seen in the GTT results of 70 patients who would have been eligible for this study. With an α of 0.05, power of 0.80, and 2-sided statistical significance testing, an estimated 16 participants would be needed to detect a medium anticipated effect size of 0.6 standard deviation (20 mg/dL lower mean 1-hour GTT result produced by the almond preload).

This study was approved by the University of Texas-Houston Institutional Review Board. Each

participant received a \$50 VISA gift card after the first GTT. If they qualified for the almond GTT, they received another \$50 gift card after completing the second test.

Data Analysis

The primary dependent variable was the mean difference in plasma glucose 1 hour after a 75-g glucose challenge with versus without the almond preload. The Student t test and area under the curve (AUC) were used to analyze the difference between standard 75-g GTT results and GTT results following the almond preload, as well as potential confounders. Statistical significance was set at P <.05 (2-sided). Food records were analyzed using the US Department of Agriculture National Nutrient Database for Standard Reference,¹³ when possible, or from the more item-specific MyFitnessPal.com database¹⁵ when foods were not clearly identifiable in the US Department of Agriculture National Database. All statistical analyses were performed with MedCalc software version 13.2.2.0 (MedCalc Software, Ostend, Belgium).

Results

Of the 53 patients who enrolled in the study, 20 met the inclusion/exclusion criteria and completed the study. Eight never did the standard GTT, 2 did the standard OGTT but not the almond GTT, and 23 did not qualify for the almond GTT. Of the nonqualifiers, 17 had a 1-hour glucose <160 mg/dL and 6 were diagnosed with diabetes. The mean interval between the 2 GTTs was 10.5 days (median, 7 days).

The mean age of the 20 study completers was 60.8 years (median, 66 years; range 18–90 years); 13 were men, 7 were women. Self-identified race/ ethnicity was non-Hispanic white in 11 completers, Hispanic in 6, Asian in 2, and African American in 1. The mean body mass index (BMI) of study completers was 29.5 kg/m² (median, 28.9 kg/m²; range, 21–38 kg/m²); 8 were obese (BMI \geq 30 kg/m²), 8 were overweight (BMI 25–29 kg/m²), and 4 were in the ideal weight range (BMI 20–24 kg/m²). The mean HbA_{1c} level was 6.0%.

The mean standard GTT results were 100.0 mg/dL fasting; 191.7 mg/dL at 1 hour; and 138.0 mg/dL at 2 hours (Table 1). Fourteen met 1 or more prediabetes diagnostic criteria; 5 had a 2-hour glucose \geq 140 mg/dL as their only prediabetes cri-

Table 1. Mean Glucose Tolerance Test Results and Potential Co	onfounders
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	Standard GTT	Almond GTT	Difference between Standard and Almond GTT, mg/dL (%)	Statistical Significance (2-Sided P Value)
Glucose (mg/dL)				
Fasting	100.0 (9.3)	97.1 (8.3)	-2.9 (2.9)	.14
1-hr	191.7 (20.6)	154.6 (37.8)	-37.1 (19.4)	<.001
2-hr	138.0 (35.4)	118.5 (27.6)	-19.5 (14.1)	.02
Carbohydrate intake (g)	228.1 (88.6)	195.1 (73.3)	-33.0 (14.5)	.18
Physical activity (minutes)				
Mean (SD)	94.6 (125.4)	117.9 (170.7)	+23.3 (24.6)	.15
Median	50	45	-5 (10)	NA
Perceived stress level	17.5 (5.6)	15.9 (4.9)	-1.6 (9.1)	.20

Data are mean (standard deviation [SD]) unless otherwise indicated. Bold values indicate statistically significant difference. GTT, glucose tolerance test; NA, not available.

terion, 4 had a fasting glucose of 100 to 125 mg/dL as their only prediabetes criterion, and 5 had both fasting glucose of 100 to 125 mg/dL and 2-hour glucose of 140 to 199 mg/dL. Six individuals had elevated 1-hour glucose \geq 160 mg/dL as their only abnormal glucose value.

Thirty minutes after almond ingestion, the baseline GTT glucose was 2.9 mg/dL lower (P = .14) than the standard GTT fasting glucose (Table 1, Figure 1). The mean 1-hour glucose was 37.1 mg/dL (19.4%) lower (P < .001) after almond ingestion than in the standard GTT. All participants' GTT values are shown in Table 2. Eight individuals showed a marked hypoglycemic effect 45 to 110 mg/dL (24% to 58%) lower and 4 showed a lesser hypoglycemic effect of 22 to 32 mg/dL (12% to 17%) for the 1-hour glucose value.





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The responses of the 4 best responders are shown in Figure 2.

The mean 2-hour glucose difference was 19.5 mg/dL (14.1%; P = .02). After almond ingestion, individual 2-hour concentrations were markedly lower by 78 to 96 mg/dL (41% to 50%) in 3, moderately lower by 20 to 39 mg/dL (10% to 20%) in 6, and not significantly different in 11 participants. Of the 8 participants who did not show a statistically significant effect at 1 hour, 2 were moderately lower (24 to 39 mg/dL) 2 hours after the almond preload.

Table 2.	Individual	Glucose To	lerance Test	Results	(mg/dL)
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	Fasting (Glucose	1-Hour	Glucose	2-Hour Glucose		
Participant	Standard	Almond	Standard	Almond	Standard	Almond	
1	92	94	178	91	106	93	
2	104	100	214	192	156	173	
3	83	86	179	87	182	86	
4	102	106	194	170	152	139	
5	122	105	178	133	115	92	
6	97	87	169	164	86	91	
7	101	97	167	174	81	120	
8	114	117	208	105	179	101	
9	94	93	166	156	131	126	
10	97	97	230	216	191	167	
11	90	99	185	190	110	109	
12	97	96	201	152	116	86	
13	93	95	173	159	141	102	
14	95	94	162	133	162	138	
15	100	105	212	102	185	100	
16	107	89	216	153	138	126	
17	117	105	212	213	182	172	
18	95	104	209	162	141	121	
19	99	87	176	144	124	119	
20	100	85	204	195	82	109	



Figure 2. Glucose tolerance test results for the best responders to the almond preload.

Almond ingestion reduced the AUC by 15.5% (P < .001). The 12 who responded positively to the almond preload lowered their 1-hour glucose by a mean of 59 mg/dL (30.2%) and lowered their 2-hour glucose by a mean of 31 mg/dL (21.8%). For the 8 nonresponders, the mean 1-hour glucose was only 8 mg/dL lower and the mean 2-hour glucose was 5 mg/dL higher after the almond preload. The patient characteristics showed no discernible predictors of good response with respect to age, sex, ethnicity, BMI, HbA_{1c}, insurance status, or family history of diabetes (Table 3). The only difference in the standard GTT results of responders versus nonresponders was a higher mean 2-hour glucose in responders (146 vs 126 mg/dL; Table 4). Nine of the 14 participants with prediabetes responded well, whereas 3 of the 6 with isolated 1-hour glucose concentrations responded well to the almond preload.

Analysis of the dietary, exercise, and stress questionnaire data for the day before the GTTs showed no significant difference in carbohydrate intake, physical activity, or perceived stress levels before the standard versus almond GTTs (Table 1).

Three study enrollees whose standard GTT showed diabetes (mean fasting glucose, 137 mg/dL; 1-hour glucose, 228 mg/dL; 2-hour glucose, 225 mg/dL) completed the almond GTT to see whether the almond preload affected more severe glucose intolerance. None of their results were significantly different after the almond preload (mean fasting glucose, 126 mg/dL; 1-hour glucose, 238 mg/dL; 2-hour glucose, 215 mg/dL).

Discussion

Ingesting ¹/₂ ounce of almonds 30 minutes before a GTT markedly lowered the 1-hour glucose in 40% of the participants and moderately lowered it in another 20%. Almond ingestion also markedly or moderately lowered the 2-hour glucose concentration in 48%, including 7 of 10 (70%) of those with prediabetic GTT 2-hour values of 140 to 199 mg/dL. The 10 participants (50%) who had 2-hour standard GTT values <140 mg/dL had relatively

Table 3.	Characteristics of	f Positive	Responders	to the A	lmond Pre	load*
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	ID	Age (years)	Sex	Race	Ethnicity	BMI (kg/m²)	HbA _{1c}	FamHxDM [†]	Insurance
Marked hypoglycemic response	2	48	Male	White	Non-Hispanic	25	5.8	0	Private
) <u>, , , , , , , , , , , , , , , , , , ,</u>	4	75	Male	AA	Non-Hispanic	33	6.3	0.5	Private
	6	71	Male	White	Non-Hispanic	29	6.3	0	Medicare
	11	46	Male	White	Non-Hispanic	37	6.1	0.5	Private
	17	78	Female	White	Hispanic	22	6.3	0	Medicare
	28	67	Female	White	Hispanic	31	6.0	1.0	Medicare
	30	90	Male	White	Non-Hispanic	23	5.3 ¹	1.5	Medicare
	40	45	Female	White	Hispanic	28	5.9	0	Medicaid
Moderate hypoglycemic response	4	67	Female	White	Non-Hispanic	30	6.4	0.63	Medicare
	14	18	Female	White	Hispanic	36	6.0	0.75	Uninsured
	19	70	Male	White	Hispanic	26	5.7	0	Medicare
	20	38	Male	White	Non-Hispanic	36	6.3	0.5	Private

*Random glucose of 145 mg/dL led to recruitment into the study, with a subsequent abnormal baseline glucose tolerance test. *Values for each relative: 1st degree (parent/sibling/child) = 0.5; 2nd degree (grandparent, grandchild, uncle, aunt) = 0.25; 3rd degree (1st cousin, great grandparent, great grandchild, great aunt or uncle) = 0.125.

(1st cousin, great grandparent, great grandennd, great aunt of uncle) = 0.125.

AA, African American; BMI, body mass index; FamHxDM, family history of diabetes.

little room for improvement in this secondary outcome variable. The mean differences between the standard GTT and the GTT after almond ingestion at 1 and 2 hours (19.4% and 14.1%) substantially exceeded test-retest variations in serial GTT values (individual coefficient of variation, 7.8%).¹⁶

The findings support the study hypothesis that "priming the pancreatic pump" with a low-calorie preload 30 minutes before GTT significantly decreases the hyperglycemic response at 1 hour in glucose-intolerant individuals who do not have diabetes. The magnitude of the AUC reduction (15.5%) was less than that produced by preloading with fructose and soya-yogurt (25% to 40%),^{8,9} but more than the results of the study by Tan and Mattes,¹² who used a larger almond snack (1.5 vs 0.5 oz) 2 hours before testing. This almond preload was more nutritionally balanced than fructose, and it contained half the calories of the soya-yogurt

Table 4. Mean Glucose Tolerance Test Results forThose Responding to Almonds versus Nonresponders

	Sta	ndard G	TT	Almond GTT				
	Fasting	1-Hour	2-Hour	Fasting	1-Hour	2-Hour		
Responders $(n = 12)$	101	194	146	99	135	115		
Nonresponders $(n = 8)$	99	188	126	95	183	125		

Data are mean values.

GTT, glucose tolerance test.

preload and one-fourth to one-half the calories in almond meal supplementation studies. $^{9,10}\,$

Shelled whole almonds can be conveniently carried in a ziplock bag. Almonds stay fresh for several months when stored at room temperature¹⁷ and can be eaten before workplace, fast food, or restaurant meals. The 30-minute premeal timing seems reasonably convenient. When bought in bulk (16 to 36 oz) a half-ounce serving of raw or roasted almonds costs 20 to 30 cents,^{18,19} so the almond appetizer is practical for families at most socioeconomic levels.

The main mechanism of action in those responding positively to the almond preload may be stimulation of release of stored insulin 30 minutes sooner than insulin release stimulated by the 75-g glucose load. The slightly lower, rather than higher, glucose concentration in the first GTT sample after almond ingestion versus the standard GTT supports this notion. Other potential mechanisms include the almonds' fiber content increasing the viscosity of intestinal contents and hindering glucose diffusion, and its fat content slowing gastric emptying time, thus delaying glucose aborption.¹²

The marked differences between the results of responders and nonresponders may be attributable to a variable combination of 2 factors. It seems likely that some of the nonresponders had more severely impaired production and/or release of insulin. The other plausible explanation is a greater degree of insulin resistance in some of the nonresponders.

Nonsignificant trends toward lower carbohydrate intake, more physical activity, and lower perceived stress the day before the almond GTT each should tend to neutralize the other's effect on the GTT results. Lower carbohydrate intake before a GTT tends to produce higher values on the test²⁰ because insulin production tends to decrease when carbohydrate intake is lower. More physical activity²¹ and lower perceived stress²² tend to produce lower GTT values.

Limitations

The study was conducted in a single clinical site. Although the study sample was fairly diverse with respect to ethnicity, socioeconomic, and insurance status, the sample size was relatively small, predominantly male, and mostly middle-aged or elderly. African Americans were underrepresented. No participant was extremely obese (BMI >40 kg/m²). It is unknown whether these findings would generalize to younger or more obese individuals.

Because the sequence of the standard and almond GTTs was not randomly varied, it is possible that the results were biased by a sequencing effect. Participants' physical activity during the GTTs was not closely monitored; bias in unobserved activity variations could have influenced the results. It would have been better to administer the stress questionnaire on the day of each GTT, rather than the day before, since perceived stress can fluctuate from day to day. The 8-ounce glass of water that was drunk only before the almond GTT and the lack of standardization of water intake during the morning hours before each GTT are potential confounders but seem unlikely to have significantly influenced the glucose results.

Plasma insulin and incretin concentrations at each of the time points in the protocol could have supported or disconfirmed the hypothesized mechanism of action. Funding constraints precluded measuring insulin or incretin concentrations.

Most blood samples were analyzed in 1 hospital laboratory, but 4 of the participants' GTT samples were analyzed by an outside commercial laboratory. Since both labs adhere to tight quality control procedures, this is unlikely to have had a significant effect on the reliability and validity of the results. The results from 3 enrollees with diabetic standard GTT results showed no hypoglycemic effect. It is possible that studying more patients with severe glucose intolerance would show a hypoglycemic effect of the almond preload for some of them. Lovejoy et al²³ found no hypoglycemic effect from supplementing the diet of 34 diabetic individuals with 57 to 113 g (2 to 4 oz) of almonds in meals and snacks each day for 4 weeks. Chen et al²⁴ found that supplementation with 60 g/day of almonds for 3 months decreased the AUC for GTT glucose by 6%.

These results raise some questions: Would a larger almond preload 30 minutes before GTT work even better? What is the optimal timing for maximizing a preload's hypoglycemic effect? What other foods would have a similar or better hypoglycemic effect? How well might the study findings translate to daily life? Will eating an almond appetizer (or similarly nutritious low-calorie snack) 30 minutes before meals lower peak postprandial glucose consistently for good GTT responders?

Conclusion

A low-calorie almond preload "appetizer" showed promise as an option for decreasing postprandial hyperglycemia in individuals with prediabetes or isolated 1-hour postprandial hyperglycemia. Further study is needed to confirm and refine the role of a premeal appetizer in the self-care of prediabetes and to investigate its potential for helping delay or avoid progression from prediabetes to diabetes.

References

- 1. Centers for Disease Control and Prevention. National diabetes statistics report: estimates of diabetes and its burden in the United States. Atlanta: U.S. Department of Health and Human Services; 2014.
- Tabak AG, Jokela M, Akbaraly TN, Brunner EJ, Kiyimaki M, Witte DR. Trajectories of glycaemia, insulin sensitivity, and insulin secretion before diagnosis of type 2 diabetes: an analysis from the Whitehall II study. Lancet 2009;373:2215–21.
- 3. Diabetes Prevention Program Research Group. 10-Year follow-up of diabetes incidence and weight loss in the Diabetes Prevention Program Outcomes Study. Lancet 2009;374:1677–86.
- Kahn SE. The importance of β-cell failure in the development and progression of type 2 diabetes. J Clin Endocrin Metab 2001;86:4047–58.
- Fantus G. Glucose toxicity. In: De Groot LJ, Chrousos G, Dungan K, et al, eds. Diabetes endotext. South Dartmouth, MA: MD Text.com, Inc.; 2000.

Available from: www.ncbi.nlm.nih.gov/books/ NBK278934/. Accessed February 23, 2016.

- 6. Gerich JE. Is reduced first-phase insulin release the earliest detectable abnormality in individuals destined to develop type 2 diabetes? Diabetes 2002; 51(Suppl 1):117–21.
- Abdul-Ghani MA, DeFonzo RA. Plasma glucose concentration and prediction of future risk of type 2 diabetes. Diabetes Care 2009;32:S194–8.
- 8. Heacock PM, Hertzler SR, Wolf BW. Fructose prefeeding reduces the glycemic response to a highglycemic index, starchy food in humans. J Nutr 2002; 132:2601–4.
- Chen MJ, Jovanovic A, Taylor R. Utilitizing the second-meal effect in type 2 diabetes: practical use of a soya-yogurt snack. Diabetes Care 2010;33:2552–4.
- Josse AR, Kendall CW, Augustin LS, Ellis PR, Jenkins DJ. Almonds and postprandial glycemia–a doseresponse study. Metabolism 2007;56:400–4.
- Jenkins DJ, Kendall CW, Marchie A, et al. Almonds decrease postprandial glycemia, insulinemia, and oxidative damage in healthy individuals. J Nutr 2006; 136:2987–92.
- Tan SY, Mattes RD. Appetitive, dietary and health effects of almonds consumed with meals or as snacks: a randomized controlled trial. Eur J Clin Nutr 2013; 67:1205–14.
- Cohen S, Janicki-Deverts D. Who's stressed? Distributions of psychological stress in the United States in probability samples from 1983, 2006 and 2009. J Appl Soc Psychol 2012;42:1320–34.
- 14. US Department of Agriculture, Agricultural Research Service, Nutrient Data Laboratory. USDA National Nutrient Database for Standard Reference, release 28. Last modified August 13, 2016. Available from: http://www.ars.usda.gov/Services/docs.htm? docid=8964. Accessed September 29, 2016.

- My Fitness Pal [homepage]. Available from: https:// www.myfitnesspal.com/. Accessed September 29, 2016.
- Kosaka K, Mizuno Y, Kuzuya T. Reproducibility of the oral glucose tolerance test and the rice-meal test in mild diabetics. Diabetes 1966;15:901–4.
- Cornacchia R, Amodio ML, Colelli G. Effects of storage temperature and duration on quality of unshelled and shelled almonds. Acta Hortic 2012;934: 783–90.
- Sam's Club. Fruit & nuts. Available from: http:// www.samsclub.com/sams/fruit-nuts/2259.cp. Accessed April 6, 2016.
- Amazon.com. Natural raw almonds (4 pound bag). Available from: http://www.amazon.com/Natural-Raw-Almonds-Pound-Bag/dp/B001ESHOA6. Accessed April 6, 2016.
- Caminhotto RDO, Lima FB. Impaired glucose tolerance in low-carbohydrate diet: maybe only a physiological state. Amer J Physiol Endocrin Metab 2013;305:E1521.
- Short KR, Pratt LV, Teague AM. The acute and residual effect of a single exercise session on meal glucose tolerance in sedentary young adults. J Nutr Metab 2012;2012:278678.
- Dungan KM, Braithwaite SS, Preiser JC. Stress hyperglycaemia. Lancet 2009;373:1798–807.
- Lovejoy JC, Most MM, Lefevre M, Greenway FL, Rood JC. Effect of diets enriched in almonds on insulin action and serum lipids in adults with normal glucose tolerance or type 2 diabetes. Am J Clin Nutr 2002;76:1000–6.
- Chen CM, Lin JF, Huang CL, et al. Effect of almonds on glucoregulation and CVD risk factors in patients with type 2 diabetes. FASEB J 2015;29(1 Suppl):912.3.