

Validation of an FFQ to assess dietary protein intake in type 2 diabetic subjects attending primary health-care services in Mali

A Coulibaly, H Turgeon O'Brien and I Galibois*

Département des Sciences des Aliments et de Nutrition, Faculté des Sciences de l'Agriculture et de l'Alimentation, Université Laval, Québec (QC), Canada G1K 7P4

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Abstract

Objective: To validate a 53-item quantitative FFQ (QFFQ) for the assessment of dietary protein intake in type 2 diabetic outpatients in Bamako, Mali.

Design: Consumption of protein-containing foods over the week preceding the interview was measured with a 7 d QFFQ and compared with intakes measured with 48-h recalls.

Setting: Centre National de Lutte contre le Diabète.

Subjects: Seventeen male and forty female adults with type 2 diabetes.

Results: Correlation between protein intakes estimated using the QFFQ and 48 h recalls was 0.63 ($P < 0.0001$). There was no significant difference between the two methods concerning the total protein daily intakes and intakes per kilogram of body weight. The QFFQ indicated that foods of animal origin were a lesser source of protein. Animal protein intake did not differ between men and women but sources did. In men, the main sources were beef (54% of total animal protein), fish (15%) and milk powder (8%). In women, the principal sources were fish (28%), beef (20%) and birds (13%). In contrast, plant protein intake was significantly higher in men than in women ($P = 0.01$), but the same plant foods contributed in similar proportions for both genders, rice being by far the greatest source (47% of plant protein in men, 53% in women).

Conclusion: The QFFQ developed in this study is a valid tool to evaluate dietary protein intakes in Malian diabetic subjects. While the total protein intakes were low in both men and women, differences in choices and amounts of protein food sources were shown.

Keywords
Type 2 diabetes
Protein intake
Food-frequency questionnaire
48 h dietary recall

Type 2 diabetes mellitus is an important health issue worldwide. Black Africans living in rural and urban regions of Africa are no exception⁽¹⁾. In Mali, the estimated number of people afflicted with diabetes was 140 000 in 2000, and it is expected to rise to 405 000 by 2030⁽²⁾. The increasing prevalence of type 2 diabetes in sub-Saharan African regions can be partly ascribed to modernisation and adoption of Western lifestyle with an associated increase of energy-dense diets, reduced physical activity and obesity⁽³⁾.

Nutrition therapy is an integral part of the treatment of diabetes mellitus and patient self-management^(4–8). Classically, more emphasis has been placed on the relative amounts and types of carbohydrate and fat to include in the diet^(5–10), but some studies have also reported beneficial effects of increasing dietary protein in type 2 diabetes^(11–13). This could be related to the fact that proteins do not increase plasma glucose concentrations in subjects with controlled diabetes, while ingested proteins are just

as potent a stimulant of insulin secretion as carbohydrates^(14–17). Thus, food proteins could contribute to an improvement in metabolic control. In the general population, the WHO/FAO recommend protein intake to account for 10–15% of the total energy⁽⁹⁾. The dietary intake of protein for individuals with diabetes and normal renal function should range between 15% and 20% of the total energy, according to the American Diabetes Association^(5,6).

In sub-Saharan Africa, particularly in Mali, it has been reported that the availability of protein per capita for the general population was 60 g/d^(18,19) in contrast with the USA where the food supply provided an average of 113 g per capita per day in 2004⁽²⁰⁾. In addition, Mali is currently affected by a strong devaluation of its currency (CFA franc). This places the population at great risk as it has reached a limit in its capacity for adaptation, which includes not only the quality but also the quantity of food consumed⁽²¹⁾. These factors may indicate that the protein

*Corresponding author: Email isabelle.galibois@aln.ulaval.ca

intake of a large number of Malian diabetic patients could be inadequate, in terms of both amount and quality.

In low-income countries, there is a lack of cost-effective dietary assessment methods⁽²²⁾; therefore it is necessary to develop quantitative methods to assess food and nutrient intake. The FFQ is currently the method most often used for assessing dietary intake in large epidemiological studies in industrialised countries⁽²²⁾. It represents a practical and cost-effective alternative to diet recalls and diet histories⁽²³⁾.

The aim of the present study was to conduct a nutritional survey in a group of type 2 diabetic patients in Mali in order to validate a 7 d FFQ developed to quantify and characterise the usual dietary protein intake.

Methods

A detailed description of the study methods has been presented elsewhere⁽²⁴⁾; therefore, only a brief summary is given here.

Subjects and ethics

The study was undertaken in a primary health-care service for diabetes: the Centre National de lutte contre le Diabète (CNLD) in Bamako, Mali.

Fifty-seven adult Malians aged between 25 and 75 years, diagnosed with type 2 diabetes and not treated with insulin, were included in the study. Participants were among the outpatients who attended primary health-care services. The patients visiting the clinic received information from their physician. In order to facilitate patients' adherence in the study, physicians were in charge of explaining the study protocol and verifying the eligibility criteria, which were checked again by a member of the research team (A.C.). The study was approved by the research ethical committee of Laval University. A written informed consent was obtained from subjects prior to their inclusion in the study.

The sample comprised more women than men (forty females *v.* seventeen males), largely reflecting the composition of the CNLD clientele as it was observed on the days of data collection. However, it could not be determined whether this is representative of the actual type 2 diabetic adult population in Mali.

Study design

This study was conducted on site over a 5-month period, between May and October 2005, corresponding to the wet season in Mali. Each subject individually met A.C. and the interviews were conducted in a local dialect, Bamana. The first interview comprised anthropometric measurements, a general questionnaire, a 7 d quantitative FFQ (QFFQ) and a dietary recall. Most subjects came back during the following weeks for another interview with A.C. to complete a second dietary recall.

Anthropometric measurements and general questionnaire

Weight was measured with light clothing on a digital scale to the nearest 0.1 kg; standing height was measured without shoes using a wooden measuring board and tape with a precision of 0.1 cm. The BMI was then calculated by dividing weight (kg) by the square of height (m).

After anthropometric measurements were taken, each participant was submitted to a general questionnaire to collect information on diabetes duration and treatment and on sociodemographic characteristics.

Dietary recalls

In the present study, participants were asked to recall their previous 48 h intake of food and beverages. It was shown that a 48 h recall is able to rank participants appropriately with respect to most nutrients and many foods, and is superior to a single 24 h recall⁽²⁵⁾. All participants completed one (*n* 14) or two (*n* 43) 48 h recalls. The nutrient content was analysed using the nutritional analysis software NUTRIFIQ developed at the Département des Sciences des Aliments et de Nutrition at Laval University in Québec, Canada, and based on the 2001 Canadian Nutrient File⁽²⁶⁾. To complete the database, nutritive values of foods from the Malian Food Composition Table⁽²⁷⁾ were added to the Canadian Nutrient File.

The 7 d quantitative FFQ

The QFFQ developed for the present study covered the amount of all protein-containing foods consumed during the 7 d preceding the interview. Using the Food Composition Table for Mali⁽²⁷⁾ and the Food Composition Tables for use in Africa⁽²⁸⁾, most food items and dishes consumed in Mali that were a source of protein (that is, supplying at least 1 g protein per 100 g) were identified. The QFFQ was pre-tested with a few diabetic patients (*n* 5) in order to ensure completeness and functionality of the questionnaire. These patients were not included in the actual study.

The QFFQ contained a list of fifty-three food items divided in two main categories: animal proteins and plant proteins. The animal protein group comprised meat, offal, poultry, fish, eggs and milk, as well as combination dishes containing these foods, while the plant protein group included dishes of beans, soya, peas, groundnut, rice, maize, sorghum, millet, wheat and tubers. Also, open-ended questions on other protein foods consumed were included at the end of the QFFQ. Food models and usual utensils were used to help participants assess the amounts eaten. In Mali, a significant proportion of daily intake is made up of foods and beverages bought from street vendors. When participants were asked to describe the size of the portions they ate, they would often refer to the price they paid. Local street foods were also purchased from vendors by A.C. in order to determine the weight of the portion by the price paid. Portion sizes of foods

recorded in household measures were also converted to weight equivalents.

For each food in the list, a protein conversion factor was derived from food composition tables. The conversion factor is a number between 0.01 and 1.00, which represents the amount of protein in 1 g of food. For each participant, the protein content of the average portion consumed on a daily basis was calculated by multiplying the weight of the portion (in g) by the conversion factor. A summation was made for the animal protein foods and the plant protein foods, as well as for the total daily protein intake.

Of the fifty-seven participants included in this study, two did not complete the FFQ; so the dietary protein intakes from the QFFQ were calculated for fifty-five participants.

Biochemical analyses

In the days following the first interview, the subjects were asked to go to a private laboratory of biomedical analyses (Laboratoire ALGI) in Bamako to undergo a blood test. Fasting venous blood samples were collected from patients using tubes containing ethylenediaminetetraacetic acid (EDTA) for glycosylated haemoglobin (HbA1c) analysis, and into tubes containing sodium fluoride for glucose analysis. Blood sample collections and the analyses were carried out directly in the private laboratory. Of the fifty-seven participants included in this study, seven failed to go to the laboratory and did not undergo the blood sampling.

Glucose levels were measured by enzymatic methods using a glucose RTU kit from Biomerieux. HbA1c was measured using the D-10 automat system of BIO-RAD, an HPLC system that operates without pre-treatment of the sample and with a restricted intervention by the user. The D-10 technique for HbA1c is certified by the 'National Glycohemoglobin Standardization Program' (NGSP), and has proven its traceability compared to the reference method of the 'Diabetes Control and Complications Trial'⁽²⁹⁾.

At each meeting with A.C. in the CNLD, capillary fasting blood glucose was also measured for each participant using a blood glucose meter and test strips from Ascencia Contour, Bayer.

Statistical analysis

Statistical analyses were performed using SAS release 9.1 (SAS Inc., Cary, NC, USA). The normality of data was tested before statistical analyses. Results are expressed as mean and *sd* unless otherwise stated. Comparisons between groups were undertaken using independent sample Student's *t*-tests. Comparisons were also made between the two methods using the paired *t*-test. Pearson's correlation coefficients were calculated to measure the association between the two dietary intake methods. Statistical significance was set at $P < 0.05$.

Results

The mean age of the participants was 54.5 (*sd* 9.4) years and their diabetes duration was 3.5 (*sd* 3.8) years. More than three-quarters of the subjects were married. In subjects' households, the meals were shared by an average of seventeen family members. About a quarter of the subjects were employed. Forty-two per cent did not have any schooling; approximately 16% went to university, while 42% of subjects had reached either a primary or a lycée/college education level.

Only one-tenth of participants took no medication and managed their diabetes only with diet, whereas one-third took medicinal plants with or without oral hypoglycaemic agents. About half of the subjects who took oral hypoglycaemic agents used drugs of the sulfonylurea class, mainly Amarel[®], Glucophage[®], Hemidaonil[®] and Daonil[®]. Regarding frequency of intake, 17.5% of the subjects took the oral hypoglycaemic agents once per day, 35% took them twice, 30% took them three times per day and finally, only 2% of subjects took these drugs four times per day. When used, the medicinal plants were mainly taken in the form of infusion. More than three-quarters of the users took them daily; only two subjects took them on a weekly basis and one only occasionally.

Concerning dietary advices, all participants, except two, said that they had received nutritional counselling related to their diabetes. Un-individualised nutritional advice given to patients at the CNLD focuses on three essential points which are: (i) foods to be avoided: sugar, banana, cream, condensed milk, cake, dried fruits, honey, fruits with syrup, fruit juice; (ii) foods to be eaten in restricted amounts, such as meat and fish, no more than 250 g/d; and starchy foods to be measured with a 400 ml bowl: white rice, steamed rice, fonio (an African cereal crop low in protein), potato, yam, millet, corn, peas, beans; and (iii) vegetables allowed without measurement. Other advice were related to lifestyle habits: 30 min of physical activity per day and weaning from smoking were recommended.

Clinical characteristics and energy and macronutrient intakes of participants according to the 48 h recalls are presented for each gender in Table 1. There was no significant difference between men and women for age and weight. BMI was larger for women than for men. On the contrary, height and daily intakes of energy were significantly higher in men than in women. However, the contribution of macronutrients expressed as a percentage of total energy was not different between men and women. Fasting blood glucose values were also similar for both genders. Although HbA1c tended to be higher in men than in women, the difference was not statistically significant.

Protein intake estimates obtained with the 48 h recalls and with the QFFQ were compared. Using Pearson's test of correlation, positive and significant correlations were

Table 1 Anthropometric characteristics, metabolic control and dietary intakes of participants according to 24 h dietary recalls (*n* 57)

Variable	Men (<i>n</i> 17)		Women (<i>n</i> 40)		<i>P</i> value
	Mean	SD	Mean	SD	
Age (years)	56.4	10.6	53.7	8.9	NS
Height (m)	1.70	0.07	1.63	0.06	0.0004
Weight (kg)	69.7	9.7	72.4	16.7	NS
BMI (kg/m ²)	24.1	3.8	27	5.3	0.043
HbA1c (%)	8.2	3.1	7.6	2.1	NS
Fasting plasma glucose (mmol/l)	7.3	3.4	7.2	3.7	NS
Capillary fasting blood glucose (mmol/l)	7.7	2.9	7.9	2.7	NS
Energy (kJ)	9792	2382	6984	2332	0.0001
Carbohydrate (% of energy)	57	9	56	8	NS
Fat (% of energy)	31	8	30	8	NS
Protein (% of energy)	12	3	12	2	NS

HbA1c, glycosylated haemoglobin.
P < 0.05.

Table 2 Daily protein intake of participants by gender according to the 48 h recalls (*n* 57) and the QFFQ (*n* 55)

Variable	48 h recalls		QFFQ		<i>P</i> value†
	Mean	SD	Mean	SD	
Daily intake					
All subjects	58	26	60	22	NS
Men‡	73	27	68	23	NS
Women§	51*	24	56	20	NS
Intake per kg body weight					
All subjects	0.8	0.4	0.9	0.4	NS
Men	1.1	0.4	1.0	0.4	NS
Women	0.7*	0.4	0.8	0.3	NS

*Significantly different from men (*P* < 0.01).

†Significance of the difference between mean recall and quantitative FFQ (QFFQ).

‡*n* 17 for dietary recall, *n* 16 for QFFQ.

§*n* 40 for dietary recall, *n* 39 for QFFQ.

found between daily protein intakes using the QFFQ and 48 h recalls (all subjects: $r = 0.63$, $P < 0.0001$; women: $r = 0.61$, $P < 0.0001$; men $r = 0.59$, $P = 0.02$). Results of the *t*-tests are shown in Table 2, where the total protein daily intakes and intakes per kilogram of body weight are also presented. There were no significant differences between the two methods.

The staple foods that were the predominant providers of protein in the diets of men and women according to the QFFQ are reported in Table 3. Concerning animal proteins, the total daily intake did not differ between men and women, but contributions of some food sources did. In men, the main sources of animal protein were in decreasing order: beef (54% of total animal protein), fish (15%) and milk powder (8%). In women, the principal sources were fish (28%), beef (20%) and birds (chicken, guinea fowl, pigeon) (13%). In absolute amounts, the daily intake of beef ($P = 0.04$) was significantly higher in men than in women, while women consumed more milk curds than men ($P = 0.01$). For plant proteins, the picture was somewhat different. Total daily intake was significantly higher in men than in women ($P = 0.01$), but the same plant foods contributed in similar proportions to

the total intake for both genders. Thus, rice was by far the largest source of plant protein (47% in men, 53% in women), followed by bread (12% in men and women) and groundnut (11% in men, 9% in women).

Discussion

In the context that information on the diet of Malian diabetic patients is scarce and considering that dietary protein could contribute to the improvement in blood glucose control, we developed an interviewer-administered QFFQ to measure the intakes of protein in the habitual diet of these subjects. The purpose of the present study was to validate this QFFQ in a group of type 2 diabetic patients attending a primary health-care service in Mali, using 48 h recalls as the reference method.

Results indicate that there were no significant differences in the intakes of protein evaluated with the QFFQ and the 48 h recalls. A significant positive correlation was found between the QFFQ and the 48 h dietary recalls for protein intake ($r = 0.63$), which was within the expected range and similar to the findings of Rodriguez *et al.*⁽³⁰⁾ ($r = 0.53$). It has been previously reported that the correlation coefficient of a nutrient should range from 0.40 to 0.70 in order to produce a good agreement between assessment methods^(31,32). Some studies on subjects without diabetes⁽³³⁾ have found a higher ($r = 0.76$), weaker or no correlation^(34,35) in dietary protein. In adults with type 1 diabetes, Riley and Blizzard⁽³⁶⁾ investigated the characteristics of a FFQ in measuring dietary intake. They have reported a weak correlation ($r = 0.38$) for dietary proteins. This difference could be due to the reference method. They used 2 d weighed dietary records, while in our study the reference method was the 48 h recall.

Although we found positive and significant correlations and no significant difference in the daily intakes between the two dietary methods, only the 48 h recalls showed significant differences between men and women in total protein intake and in intake per kilogram of body weight.

Table 3 Main sources of animal and plant protein in diets of type 2 diabetic men and women in Mali

Source	Amount (g/d)					Protein (g/d)				
	Men (n 16)		Women (n 39)		P value	Men (n 16)		Women (n 39)		P value
	Mean	SD	Mean	SD		Mean	SD	Mean	SD	
Animal protein										
Beef	48.2	54.6	17.2	21.4	0.04	13.0	14.7	4.6	5.8	0.04
Mutton	4.4	11.0	3.7	9.3	0.82	1.2	3.0	1.0	2.5	0.82
Goat	0.5	2.1	1.9	11.4	0.46	0.14	0.6	0.5	3.1	0.46
Offal	1.2	3.4	2.6	6.6	0.52	0.3	0.9	0.6	1.4	0.27
Bird	5.1	12.0	11.1	27.8	0.26	1.4	3.2	3.0	7.5	0.26
Fish	13.3	10.4	24.6	38.1	0.09	3.6	2.8	6.6	10.3	0.09
Eggs	0.1 ^a	0.4	0.1 ^a	0.2	0.99	1.2	3.1	1.2	1.8	0.99
Milk powder	7.5	14.3	10.0	13.1	0.53	2.0	3.7	2.6	3.4	0.53
Liquid milk	21.9	42.0	52.4	114.5	0.15	0.9	1.7	2.1	4.7	0.15
Milk curds	0.0	0.0	18.9	45.0	0.01	0.0	0.0	0.7	1.8	0.01
Total						24	18	23	16	NS
Plant protein										
Beans	22.3	43.3	17.0	37.8	0.65	1.6	3.0	1.3	2.8	0.73
Groundnut	24.3	26.7	17.4	24.9	0.36	4.9	5.9	3.0	3.9	0.25
Peanut sauce	41.7	57.0	21.5	25.6	0.19	1.2	1.7	0.6	0.8	0.19
Rice	700.7	336.1	594.3	318.8	0.27	20.6	9.4	17.5	9.6	0.28
Maize	134.5	224.5	49.0	110.8	0.16	2.9	4.7	1.1	2.2	0.14
Sorghum	48.2	178.2	125.5	237.7	0.25	1.1	3.8	2.9	5.3	0.22
Millet	150.4	289.1	59.1	148.1	0.24	3.3	6.8	1.3	3.3	0.27
Macaroni	72.8	107.8	33.1	62.3	0.18	1.4	2.1	0.7	1.2	0.18
Bread	74.8	61.1	56.0	57.6	0.28	5.2	4.2	3.9	4.0	0.28
Yam with meat	30.3	53.2	22.7	45.8	0.59	0.3	0.5	0.2	0.4	0.59
Boiled yam	8.5	23.2	5.3	13.5	0.61	0.4	1.2	0.3	0.7	0.61
Cassava	18.7	64.0	4.1	10.1	0.38	0.7	2.6	0.2	0.4	0.38
Potato	12.0	26.0	16.3	33.6	0.65	0.2	0.5	0.3	0.7	0.65
Sweet potato	6.6	24.9	1.1	4.3	0.4	0.1	0.2	0.01	0.04	0.4
Total						44	16	33	14	0.01

^aNumber of eggs consumed per day.

This might be due to under- and overestimation of intakes, depending on the dietary assessment method. It has been shown that FFQ can both under- and overestimate the intakes of specific nutrients⁽³⁴⁾. In fact, many validation studies have reported that FFQ, when compared to food records or 24 h recalls, overestimate nutrient intakes^(22,23,30,33,37). In contrast, other studies have reported that FFQ did not systematically overestimate nutrient intakes^(34,38–40).

Regardless of dietary methods, the present study showed that dietary protein intakes in a group of type 2 diabetic men and women in Mali were similar to what is observed for protein availability (around 60 g per capita per day) in the diet of the general Malian population^(18,19). They were also quite similar to the dietary protein intakes of South African black men and women with type 2 diabetes (63 and 50 g/d, respectively)⁽⁴¹⁾, but lower than that of Ghanaian type 2 diabetics (81 g/d)⁽⁴²⁾. Compared to the protein availability of the US population (113 g per capita per day) in 2004⁽²⁰⁾, the lower protein intake of the participants in this study could be due to the reduced financial means and to the large number of family members, which could limit the access to excellent sources of protein such as meat and fish. In fact, Torheim *et al.*⁽²²⁾ have reported in their study conducted in a Malian village that meat and fish were rarely eaten.

The fact that food patterns in Mali are seasonal^(19,22) could explain the low intake of protein by the diabetic patients. Indeed, our study was conducted during the wet season, which is not the period of harvest, and hence staple foods such as cereals were not abundant. The low protein intake could also be due to the fact that patients were sometimes given conflicting advice with respect to the type of foods they were allowed to eat, and they generally appeared to have little understanding of portion sizes. We found that the main provider of proteins in their diet was plant protein, which was also shown in other studies^(18,19). But at the same time, as doctors and nurses had counselled them to eat less food that was rich in starch, our subjects may have under-reported their plant food intake to show their adherence to nutritional counselling. Although dietary carbohydrate is the major contributor to postprandial glucose concentration, it is an important source of energy, water-soluble vitamins and minerals, and fibre^(5–7). Hence, low-carbohydrate diets are not recommended in the management of diabetes. The amount of carbohydrate ingested is usually the primary determinant of postprandial response, but the type of carbohydrate also affects this response^(5–7). As carbohydrate-containing plant foods are the main provider of dietary protein in Malian diabetic patients, they should be advised to give preference to plant foods such as legumes

that could provide more protein than rice or maize, and that could also provide more fibre, which is beneficial in type 2 diabetes^(5–7). Moreover, an experimental study in subjects with type 2 diabetes reported that compared to potatoes, dried peas induced a delayed and smaller increase in postprandial plasma glucose, supporting the suggestion that type 2 diabetic patients should increase their consumption of low-glycaemic, high-fibre foods at the expense of high-glycaemic, low-fibre foods⁽⁴³⁾.

We should also keep in mind that Mali is one of the poorest countries in Africa and with the 1994 devaluation of the currency, a great impoverishment of the population was observed. The continuing impoverishment of the Malian population, the impact of large family sizes and food consumption units, and the cost of diabetes treatment pose a major challenge to patients who cannot afford more than staple foods, which are mainly plant food.

The higher consumption of dietary protein and beef in men compared to women can be explained by socio-cultural conditions prevailing in Mali such as polygamy, number of family members and the women's position in the family. In fact, women are generally housewives, with men being the main financial providers. This leaves women in a lower economic situation. Men have privileged access to food and, as seen here, have more beef in their diet than women. Moreover, women must at first satisfy their husband and children's dietary intake in order to be well regarded by their family. The presence of street food consumption in Mali^(21,44) could also explain the higher consumption of animal protein and beef in men. The fact that men have financial resources could allow them to buy street foods rich in protein such as meat, fish and eggs to supplement their dietary protein intake.

In patients with type 2 diabetes, it was reported that the simultaneous ingestion of glucose with protein in test meals significantly decreased the glycaemic response, as compared with glucose taken alone⁽⁴⁵⁾. Other single-meal studies confirmed that protein foods have a modest impact on blood glucose but a significant effect on insulin secretion^(14,15) in type 2 diabetic subjects. In addition, clinical studies have shown that an increase in dietary protein improved the metabolic control in type 2 diabetes, albeit in the context of a weight-loss diet⁽¹¹⁾ or without weight loss⁽⁴⁶⁾.

Considering these potential beneficial effects of dietary protein in type 2 diabetes, future studies should be conducted to evaluate the effect of a nutritional intervention in Malian type 2 diabetic patients without nephropathy that would aim to improve their protein intake. Also, as dietary habits depend on the time of the year, nutrition surveys should be repeated in other seasons and with larger and more representative samples to confirm the assessment of usual protein intake with the QFFQ. Finally, dietary protein intake could be validated using biomarkers such as urinary nitrogen.

Conclusion

In conclusion, although our study presents some limitations, we found that the QFFQ developed in the present study is a valid and useful tool to estimate the average daily protein intakes of type 2 diabetic patients in Mali. We also found that diabetic patients need to receive more nutrition counselling and many patients need to increase their protein intake, particularly women. However, it will be necessary to conduct nutritional surveys with larger samples to verify the actual intake of the type 2 diabetic population in Mali.

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