Impact of a Remote Monitoring Programme Including Lifestyle Education Software in Type 2 Diabetes: Results of the Educ@dom Randomised Multicentre Study

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ABSTRACT

Introduction: Telemonitoring in type 2 diabetes (T2D) is mainly based on glucose monitoring. A new type of connected device which routinely gathers data on weight, physical activity and food intake could improve patients’ diabetes control. The main aim of this study was to assess the efficacy of an at-home interventional programme incorporating such devices and lifestyle education software on diabetes control, i.e., change in HbA1c, compared to standard care.
Methods: This multicentre study randomly assigned 282 people with T2D to either a telemonitoring group (TMG) or a control group (CG) for a 1-year intervention period. While routine follow-up was maintained in the CG, TMG subjects were provided with interactive lifestyle educational software (with artificial intelligence algorithms) and connected objects (blood glucose meters, scales and actimeters) for use in their own homes and were remotely monitored by their diabetologists. Changes in HbA1c were compared between groups using a mixed linear model.

Results: The mean HbA1c dropped from 7.8 ± 0.8% (62 mmol/mol) to 7.4 ± 1.0% (57 mmol/mol) in the TMG and from 7.8 ± 0.8% (62 mmol/mol) to 7.6 ± 1.0% (60 mmol/mol) in the CG, resulting in an intergroup difference of −0.16 (p = 0.06) in favour of TMG, after adjustment for confounding factors. Within TMG, the decrease in HbA1c was greater in frequent users: −0.23% (p = 0.03) in the case of connections to telemonitoring synthesis above the median and −0.21% (p = 0.05) in the case of connections to tele-education software above the median compared to the CG. Significant weight loss was observed in the TMG but only in women (p = 0.01).

Findings: The EDUC@DOM telemonitoring and tele-education device did not highlight a significant decrease in HbA1c levels compared to routine management although a slight, albeit significant improvement in glycaemic control was observed in the frequent user subgroup as well as significant weight loss but only in women. A high level of satisfaction with the connected device was recorded among all participants.

Trial Registration: This trial was registered in the Clinical Trials Database on September 27, 2013, under no. NCT01955031 and bears ID-RCB number 2013-A00391-44.

Keywords: Type 2 diabetes; Lifestyle management; Glucose control; Telemonitoring; Tele-education; Health-related objects

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Introduction

According to current guidelines, first-line intervention in individuals living with type 2 diabetes (T2D) is based on lifestyle changes aimed at a balanced diet and frequent physical activity in order to achieve better glycaemic control and, if necessary, weight reduction [1]. Such lifestyle modifications need to be sustained with a long-term perspective, regardless of the combined pharmacological treatment regimens. Although many studies have demonstrated the ability of lifestyle interventions to improve glycaemic control in T2D, this
beneficial metabolic effect is difficult to sustain long term [2–4]. Technological innovations, including the use of health-related objects, currently make it possible to remotely monitor individuals with diabetes in their own homes. Close intervention in their daily lives is thus feasible, both as a support towards improving lifestyle and adapting treatment [5–20].

This randomised study was based on the hypothesis that tele-education and telemonitoring improve lifestyle and quality of glycaemic control in patients with T2D while rationalising the need for face-to-face consultations. The long-term goal is to reduce the incidence of diabetic complications and costs. We designed and developed Nutri-Educ, a new at-home education software, in order to support changes in lifestyle. Nutri-Educ is the only software including artificial intelligence algorithms, which allow it to deliver personalised advice to correct errors in meal composition. This software has been combined with the use of connected objects to offer an integrated telemonitoring programme [21]. Changes in HbA1c, weight and waist circumference have been reviewed over a 1-year period in T2D subjects equipped with this innovative telemonitoring programme compared to standard care. The main aim of this study was to assess the efficacy of the telemonitoring programme in improving glycaemic control, i.e., decreasing HbA1c levels compared to standard care.

METHOD

Trial Design

This open-label, randomised, controlled trial with two parallel arms, telemonitoring group (TMG) and control group (CG), involved 16 investigation sites: 5 university teaching hospitals, 1 non-teaching hospital centre, a diabetes care network linking 8 public and private health establishments and 2 private practice physicians.

Participants

The subjects enrolled in the study were aged 18 years or above, had a documented medical history of T2D, with or without insulin treatment, and with a recent HbA1c value > 6.5% (48 mmol/mol) and ≤ 10% (86 mmol/mol). They had to have an active internet connection at home. People with severe or active comorbidities, reduced mobility, serious eating disorders or who had undergone bariatric surgery were excluded [21].

Interventions

The home telemonitoring device combined biomedical data sensors (scale with impedancemetry, actimeter and blood glucose meter) with educational software [22, 23] available on tablets. Three tele-educational software programmes were given to participants in the TMG: Nutri-Kiosk—a set of self-learning quizzes for formative assessment based on nutritional knowledge; Acti-Kiosk—support in starting physical activity (assessment, information and videos); Nutri-Educ—a personalised nutritional educational software application that meets international clinical recommendations [24]. It was developed using artificial intelligence algorithms and has already demonstrated its beneficial impact in obesity and T2D patients [25–27]. Nutri-Educ was designed to help patients in the TMG group to improve their nutritional balance. The patients had to enter meal composition details into the software, which subsequently analysed the meals and proposed corrections taking into account the patients' individual profiles, dietary habits and tastes. A more detailed description of the device has already been published [21].

The monitoring interface linking TMG participants with investigators was designed as a secure web platform receiving all data from connected objects. TMG subjects sent these data to the platform on a weekly basis and had free access to this information and to their monitoring summary. The frequency of data consultation by investigators was left to their discretion. However, according to the protocol,
they were sent an email every month to encourage them to carry out the telemonitoring. A summary of all data recorded over the last 30 days was available for rapid interpretation of results in order to facilitate follow-up and the telemonitoring process. A break-down of the monthly changes also provided an overview of the progress in the monitored parameters. Modifiable alerts notified investigating physicians about certain events including hypoglycaemia, hyperglycaemic tendencies and significant weight gain or loss. Investigators had access to patient records to personalise the follow-up by modulating blood glucose, body weight and physical activity targets to be achieved over time as well as alert levels according to each participant’s profile. A secured messaging system allowed repeated and interactive discussions with telemonitored individuals to advise, encourage and coach them. Investigating physicians were required to provide a report on telemonitoring and the actions implemented in order to validate the telemonitoring process and ensure traceability of the action taken. General practitioners were given regular telemonitoring progress reports.

Telemonitoring follow-up for the TMG was planned over a 12-month period, with no face-to-face consultation scheduled in advance. Follow-up in the CG included face-to-face consultations in accordance with standard practices.

Outcomes

The primary endpoint was the intergroup change in glycated haemoglobin levels (HbA1c) over the 12-month intervention period. Changes in other clinical parameters (body weight, BMI and waist circumference) were also considered and compared between groups in the overall population and then taking gender into account. Compliance with the device, including frequency of use and its impact, was assessed. Finally, the satisfaction of individuals with diabetes and caregivers was also recorded using an ad hoc questionnaire dedicated to the device in question.

Randomisation

The individuals with T2D enrolled in the trial were randomly assigned to two groups (telemonitoring group [TMG] or control group [CG]) balanced by random block sizes and stratified by HbA1c level at inclusion [threshold: 7.5% (58 mmol/mol)], which was a priori expected to be the most relevant factor associated with changes in HbA1c levels during intervention.

Sample Size

The sample size was initially determined to detect a 0.5 mmol/mol difference in the primary efficacy endpoint over 12 months, with a power of 80% and a 5% type I error (2-tailed test). We applied the Frison and Pocock approach [28], based on a comparison of the mean of four repeated measurements between the two intervention arms. A sample size of 234 (117 per arm) was necessary, plus 20% to avoid loss of power due to loss of follow-up, resulting in a total of 282 subjects to be recruited (141 per arm).

Statistical Analysis

An intention-to-treat effectiveness analysis was carried out. Since the analysis was performed on complete data (no imputation for missing data), the intention-to-treat population comprised all randomised participants having at least one HbA1c value collected during follow-up (primary endpoint) and having no missing data regarding the confounding factors defined a priori. The proportion of subjects randomised but not analysed was < 10%.

Concerning the primary efficacy endpoint, we compared the two groups using a linear mixed-effects model for repeated measures
(MMRM), modelling all HbA1c measurements performed during the 1-year intervention period. To ensure the comparability of the two groups (TMG versus CG) and to enhance statistical power, the model was adjusted for the confounding factors defined a priori (baseline HbA1c, age, gender, diabetes complications, obesity, duration of diabetes and insulin treatment). The search for an interaction between the time of measurement and the intervention group was carried out as well as a centre effect. Additional analyses taking patient compliance (number of connections) into account were also performed. For this purpose, we considered the number of connections to data synthesis and to Nutri-Educ software, respectively. These variables were categorised into two groups based on the median value. We therefore classified the most assiduous users into TMGs+ group (number of connections to the synthesis above the median) and TMGn+ group (number of connections to the Nutri-Educ software above the median).

Changes in body weight, BMI and waist circumference during follow-up were compared using the same procedure. The analyses were also stratified by gender to assess a potential change in terms of effect.

Since the primary and secondary analyses suggested an effect dependent on compliance with the device, the correlation between the changes in HbA1c and BMI was tested according to the frequency of connections to the device (synthesis and Nutri-Educ) in the TMG.

Data analysis was performed using Stata software (Statistical Software: Release 14.0. Stata Corporation, College Station, TX).

Ethics Approval and Consent to Participate

All T2D patients participating in this research received verbal and written information about the study and were given the opportunity to ask any questions to help them understand the study. They signed a voluntary informed consent form before the research began. This study was submitted to the South-West and French Overseas Territories’ Research Ethics Committee (REC) [Comité de Protection des Personnes (CPP)] and received a favourable opinion from the REC on 27/05/2013.

RESULTS

Patient Disposition and Baseline Characteristics

Two hundred eighty-two T2D patients were enrolled in the study according to baseline HbA1c level stratification, i.e., HbA1c < 7.5% (58 mmol/mol) (n = 131) or HbA1c ≥ 7.5% (n = 151). They were then randomly assigned to two groups of 141 participants for either telemonitoring (TMG) or standard (CG) management. Nineteen participants were lost to follow-up (13 in the TMG and six in the CG) (Fig. 1).

The analysis set comprised 263 individuals (128 in the TMG and 135 in the CG) who had at least one HbA1c measurement during the follow-up period. Multivariate analyses were based on 256 participants (126 in the TMG and 130 in the CG) with no missing data among the variables entered in the model (data and confounding factors defined a priori). Finally, 242 individuals presented with an HbA1c measurement at 12 months, i.e., at the end of the intervention period.

The baseline characteristics of the study population show no difference between the two groups (Table 1).

Frequency of Device Use in Telemonitored Patients

In the TMG, the mean number of connections to the device by patient was 104 ± 78 (median value: 86) over the 12-month follow-up period, which corresponds to a frequency of about twice a week. Users could access the different device functions at each connection. Mean data synthesis (TMGs) and Nutri-Educ software (TMGn) access figures were 44 ± 49 (median value: 29) and 48 ± 61 (median value: 31), respectively, demonstrating almost weekly use. On average, TMG patients sent 14 ± 13 messages (median value: 11) to the investigators,
i.e., about one message per month. The mean messaging frequency from the investigators to the participants was 5 ± 5 (median value: 3).

**Primary Outcome: Changes in HbA1c over the Intervention Period**

During the 12-month intervention period, the mean HbA1c level dropped from 7.8 ± 0.8% (62 mmol/mol) to 7.4 ± 1.0% (57 mmol/mol) in the TMG and from 7.8 ± 0.8% (62 mmol/mol) to 7.6 ± 1.0% (60 mmol/mol) in the CG. Among individuals with measurements available at 12 months, no significant decrease in the mean HbA1c level was recorded in the TMG compared to the CG: −0.19% (95% CI [−0.43; 0.05], p = 0.12). Moreover, the remaining difference between HbA1c value achieved at 12 months and the target initially set with the investigator (available for 241 participants) was not significantly lower in the TMG compared to the CG (−0.13%, [−0.37; 0.10], p = 0.27).

After adjustment for confounding factors, the intergroup difference was −0.16% (95% CI [−0.32; 0.01], p = 0.06), in favour of TMG (Fig. 2). The change predicted by the adjusted model showed an improvement in HbA1c level from T0 to T12: 7.75% (95% CI [7.65; 7.85]) (61 mmol/mol) to 7.49% (95% CI [7.31; 7.66]) (58 mmol/mol) in the TMG and to 7.64% (95% CI [7.47; 7.82]) (60 mmol/mol) in the CG.

Taking into account stratification on baseline HbA1c, the intergroup difference in terms of change was −0.13 (95% CI [−0.32; 0.05], p = 0.16) in the lower stratum (HbA1c < 7.5%, n = 120), and −0.19 (95% CI [−0.45; 0.07], p = 0.16) in the upper stratum (HbA1c ≥ 7.5%, n = 136). We found no interaction between baseline HbA1c stratum and telemonitoring intervention regarding changes in HbA1c levels (p = 0.55).

Individuals characterised as TMGs+ (connections to telemonitoring synthesis above the median) exhibited a greater decrease in HbA1c compared to the CG: −0.23% (95% CI [−0.43; −0.03], p = 0.03), with a 12-month predicted change of −0.31% (95% CI [−0.53; −0.10]) in TMGs+ versus −0.08% [−0.26; 0.09] in CG, moving from 7.75% (61 mmol/mol) to 7.42% (58 mmol/mol) for TMGs+ versus 7.64% (60 mmol/mol) for CG. Similarly, a fall in HbA1c levels was more important in TMGn+ (connections to Nutri-Educ above the median) than in CG participants: −0.21% (95% CI [−0.41; −0.00], p = 0.05), with a 12-month predicted decrease of −0.29% (95% CI [−0.51; −0.07]) in TMGn+ versus −0.09% [−0.26; 0.09] in CG, moving from 7.75% (61 mmol/mol) to 7.44% (58 mmol/mol) for TMGn+ versus 7.64% (60 mmol/mol) for GC (Table 2).

**Secondary Outcomes: Changes in BMI and Waist Circumference During the Intervention Period**

Among individuals with data available over 12 months (n = 240), BMI decreased by 0.30 ± 1.9 kg/m² + in the TMG and increased by 0.06 ± 1.3 kg/m² in the CG resulting in a non-significant intergroup difference of −0.36 kg/m² (95% CI [−0.76; 0.05], p = 0.08) in favour of the TMG. After adjustment for confounding factors (n = 236), the average predicted difference between the two groups was −0.30 kg/m², (95% CI [−0.70; 0.09], p = 0.14) with an expected 12-month change of −0.24 kg/m² in the TMG versus 0.06 kg/m² in the CG. The expected BMI changes between baseline and the end of intervention were −0.40 kg/m² (−0.80; 0.00) in TMGs+ patients (versus 0.06 kg/m² [−0.24; 0.35] in CG patients), and −0.53 kg/m² [−0.93; −0.14] in TMGn+ patients, reflecting an anticipated respective difference of −0.46 kg/m² (−0.94; 0.02), p = 0.06 and −0.57 kg/m² (−1.05; −0.09), p = 0.02 relative to CG patients.

In patients with available waist circumference values, unadjusted (n = 206) and adjusted (n = 202) intergroup differences were −1.67 cm ([−3.27; −0.06], p = 0.04) and −1.29 cm ([−2.87; 0.30], p = 0.11), respectively, both in favour of the TMG. Intergroup differences were more pronounced in frequent users: −2.00 cm ([−3.90; −0.10], p = 0.04) between TMGs+ and CG patients and −2.64 cm
Predicted differences in HbA1c (primary end-point) were $-0.21\%$ ($[-0.48; 0.05], p = 0.12$) in women and $-0.15\%$ ($[-0.35; 0.05], p = 0.15$) in men, in favour of TMG patients.

Intergroup differences in BMI (TMG versus CG) were $-1.01$ kg/m$^2$ ($[-1.79; -0.23], p = 0.01$) in women and $0.16$ kg/m$^2$ ($[-0.27; 0.60], p = 0.46$) in men. According to the frequency of use, BMI differences of $-1.23$ kg/m$^2$ ($[-2.16; -0.31], p = 0.009$) were documented in TMGs+ women and $-1.40$ kg/m$^2$ ($[-2.31; -0.48], p = 0.003$) in TMGn+ women. In terms of body weight, women in the TMG lost $2.7$ kg ($[-4.79; -0.57], p = 0.01$) more than those in the CG group and up to $3.7$ kg ($[-6.30; -1.11], p = 0.005$) in TMGs+ and $-3.78$ kg ($[-6.25; -1.31], p = 0.003$) in TMGn+ women. No significant changes in waist circumference were observed (data not shown).

HbA1c significantly decreased over the 12-month intervention period as the number of connections to telemonitoring syntheses increased ($-0.16$ $[-0.32; -0.003], p = 0.04$). No significant correlation was found between this frequency of use indicator and change in BMI ($p = 0.28$).

The number of connections to Nutri-Educ tended to be correlated with changes in HbA1c ($-0.15$ $[-0.31; 0.02], p = 0.08$) and was significantly correlated with changes in BMI ($-0.30$ $[-0.58; -0.02], p = 0.03$).

**Device User Satisfaction (Fig. 4)**

At the end of the 12-month intervention period, 91.0% of telemonitored individuals completed the satisfaction questionnaire; 97.4% were completely satisfied or rather satisfied with device use and telemonitoring data synthesis. Very high levels of satisfaction (very satisfied or rather satisfied) were also recorded regarding the assistance provided by the device in terms of

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**Correlation Between Device Use and Changes in HbA1c and BMI (Fig. 3)**

HbA1c significantly decreased over the 12-month intervention period as the number of connections to telemonitoring syntheses increased ($-0.16$ $[-0.32; -0.003], p = 0.04$). No significant correlation was found between this frequency of use indicator and change in BMI ($p = 0.28$).

The number of connections to Nutri-Educ tended to be correlated with changes in HbA1c ($-0.15$ $[-0.31; 0.02], p = 0.08$) and was significantly correlated with changes in BMI ($-0.30$ $[-0.58; -0.02], p = 0.03$).
|                          | Total (n = 263) | CG (n = 135) | TMG (n = 128) |
|--------------------------|----------------|-------------|-------------|
| **Gender**               |                |             |             |
| Male                     | 166            | 63.1%       | 84          | 62.2% | 82          | 64.1%        |
| Female                   | 97             | 36.9%       | 51          | 37.8% | 46          | 35.9%        |
| **Age (years)**          |                |             |             |
| Median (IIQ)             | 60.6           | 13.0        | 60.3        | 14.0  | 61.3        | 12.4         |
| Mean (SD)                | 59.5           | 9.6         | 59.3        | 10.0  | 59.8        | 9.2          |
| **Randomisation stratum**|                |             |             |
| Initial HbA1c < 7.5%     | 123            | 46.8%       | 63          | 46.7% | 60          | 46.9%        |
| Initial HbA1c ≥ 7.5%     | 140            | 53.2%       | 72          | 53.3% | 68          | 53.1%        |
| **Initial HbA1c**        |                |             |             |
| Median and IIQ           | 7.6            | 1.1         | 7.7         | 1.1   | 7.6         | 1.2          |
| Mean and standard deviation | 7.8          | 0.8         | 7.8         | 0.8   | 7.8         | 0.8          |
| **Body weight (kg)**     |                |             |             |
| Median and IIQ           | 91.0           | 21.0        | 89.0        | 24.0  | 91.5        | 18.5         |
| Mean and standard deviation | 91.7          | 16.7        | 90.0        | 16.0  | 93.5        | 17.3         |
| **BMI (kg/m²)**          |                |             |             |
| Median and IIQ           | 32.2           | 5.7         | 30.8        | 5.8   | 31.6        | 6.0          |
| Mean and standard deviation | 31.8          | 5.3         | 31.2        | 5.2   | 32.5        | 5.4          |
| **Obesity (BMI ≥ 30 kg/m²)** |            |             |             |
| No                       | 99             | 37.8%       | 58          | 43.6% | 41          | 32.0%        |
| Yes                      | 162            | 62.2%       | 75          | 56.4% | 87          | 68.0%        |
| **Waist circumference**  |                |             |             |
| Mean (SD)                | 110.1          | 12.5        | 107.9       | 12.4  | 112.4       | 12.3         |
| **Duration of diabetes** |                |             |             |
| Less than 5 years        | 31             | 11.9%       | 16          | 12.0% | 15          | 11.8%        |
| Between 5 and 15 years   | 127            | 48.9%       | 65          | 48.9% | 62          | 48.8%        |
| More than 15 years       | 102            | 39.2%       | 52          | 39.1% | 50          | 39.4%        |
| **Diabetes complications**|              |             |             |
| No                       | 121            | 46.5%       | 60          | 45.1% | 61          | 48.0%        |
| Yes                      | 139            | 53.5%       | 73          | 54.9% | 66          | 52.0%        |
| **Details of complications** |            |             |             |
| Retinopathy              | 41             | 15.9%       | 25          | 19.2% | 16          | 12.5%        |
| Nephropathy              | 66             | 25.9%       | 32          | 24.8% | 34          | 27.0%        |
improving their knowledge on diabetes (87.6%) and nutrition (84.3%). Device-generated assistance was deemed positive by 88.2% of subjects for the daily management of diabetes, by 81% for ensuring a balanced diet and by 85.5% for the practice of physical activity.

Fifty-five percentage of the investigators completed the satisfaction questionnaire; 85% of them reported having completely integrated the web application functions and over 80% found it easy to very easy to use, in terms of both patient records and telemonitoring synthesis reports. Finally, 82.3% were keen to continue using the device.

**DISCUSSION**

This randomised study shows that, compared to standard management including face-to-face outpatient visits, use of the EDUC@DOM telemonitoring and tele-education device for a period of 1 year did not significantly improve glycaemic control in people with type 2 diabetes (non-significant decrease in HbA1c levels, \( p = 0.06 \)). However, the metabolic benefit was logically correlated with the frequency of use of the device, which also resulted in weight loss and a reduction in waist circumference in women. It should be noted that individuals with T2D who participated in this study had all previously benefited from conventional education programmes and were all followed up by a diabetologist prior to inclusion, which could have minimised the overall impact of this intervention. The effectiveness of such connected programmes could thus certainly be increased in patients less experienced in the management of their disease.

The growing prevalence of T2D along with the limited number of physicians in most countries makes it difficult to set up regular medical follow-up, sometimes requiring face-to-face visits, especially when glycaemic control is insufficient. The recent development of telemonitoring devices and connected objects should change the way in which diabetes is managed and allow greater flexibility in terms of follow-up.

Nutritional intervention is effective in improving glycaemic control in T2D patients but it requires significant educational and therapeutic support from caregivers. In this specific field, advances in remote communication technologies and the development of

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**Table 1 continued**

|                         | Total (\( n = 263 \)) | CG (\( n = 135 \)) | TMG (\( n = 128 \)) |
|-------------------------|------------------------|--------------------|---------------------|
| Peripheral neuropathy   | 43 (16.7%)             | 29 (22.1%)         | 14 (11.0%)          |
| Vegetative neuropathy   | 10 (4.0%)              | 6 (4.7%)           | 4 (3.2%)            |
| Coronary insufficiency  | 47 (18.4%)             | 19 (14.7%)         | 28 (22.2%)          |
| Cerebrovascular insufficiency | 12 (4.7%) | 8 (6.1%)           | 4 (3.2%)            |
| Lower limb artery disease | 25 (9.8%)        | 9 (6.9%)           | 16 (12.9%)          |
| Chronic wound           | 8 (3.1%)               | 4 (3.0)            | 4 (3.2%)            |
| Receiving insulin       |                        |                    |                     |
| No                      | 79 (30.0%)             | 42 (31.1%)         | 37 (28.9%)          |
| Yes                     | 184 (70.0%)            | 93 (68.9%)         | 91 (71.1%)          |

Number and percentage, unless otherwise stated
applications for the automated assessment of nutritional intake have also opened up new avenues. However, the hypothesis that nutritional intervention based on telemedicine and artificial intelligence (AI) solutions could be as effective as or even more effective than traditional face-to-face consultations has yet to be demonstrated.

To date, studies evaluating telemonitoring devices dedicated to diabetes have generated mixed results in terms of both lifestyle and metabolic balance [18, 29–33]. By incorporating connected objects and tele-education tools including AI algorithms, the originality of our device is that it offers personalised support for lifestyle changes, particularly regarding nutrition.

Compliance with the use of the various connected objects and educational software is undoubtedly vital in ensuring the efficacy of this care method. Indeed, the benefit observed in terms of HbA1c changes as well as changes in BMI in women depends on the frequency of connection to the educational software and patient access to telemonitoring data synthesis. In this context, it is important to emphasise the fact that, with a mean age of 60 years and rather persistent diabetes (of more than 5 years’ duration in 88% of cases), our study participants were definitely not in the highly tech-savvy population.

In our study, we chose to include T2D patients regardless of their initial level of glycaemic control. Stratified analysis according to the baseline HbA1c value did not highlight a significant difference between the two strata in terms of changes in HbA1c over the 12-month follow-up period, although a greater drop could have been expected in less well-controlled individuals, as usually reported [34, 35]. However, from a public health perspective and to avoid long-term complications [36], we initially considered that, in addition to improving glycaemic control in individuals with high HbA1c levels, maintaining subjects with slightly unbalanced diabetes at satisfactory HbA1c levels is also valuable.

Weight and BMI monitoring also shows benefits directly related to the frequency of device use although this beneficial impact appears to be more significant in women than in men. A reduction in waist circumference is also correlated with device use but no gender effect is apparent. Such positive effects on body weight and fat distribution may have been associated with improved insulin sensitivity and a reduction in cardiovascular risk factors, but these parameters were not documented in our study [37].

In T2D, blood glucose monitoring must be linked to lifestyle interventions in order to improve metabolic control [32]. Therefore, in our telemonitoring and coaching approach, we included not only a glucose meter but also other connected objects and tele-education modules targeting diet and physical activity. It is important to note that the Nutri-Educ software included in our device has already demonstrated its effectiveness in improving dietary balance, weight and glycaemic control in individuals with diabetes or obesity [23, 24].

Long-term lifestyle changes remain a key challenge in the management of T2D to eradicate often deeply ingrained habits. To this end, telemonitoring and tele-education approaches offer innovative solutions through the provision of personalised and on-going advice to modify diet and physical activity practices while respecting patients’ tastes and preferences. Such devices provide physicians and their patients with regular updates on assessments of the patients’ lifestyle in relation to the jointly set targets. In addition, AI modules are now able to drive the solutions most suited to each patient’s clinical profile. In addition to the data collected, the connected objects per se also play a
pedagogical role by boosting motivation, but
their impact is enhanced by the coaching effect
in particular [31]. Finally, remote monitoring
over several months prompts closer support and
helps maintain the desired behavioural changes
to control the disease. In line with this last
point, most users were satisfied or very satisfied
with the EDUC@DOM tool for assistance in
diabetes management and especially in lifestyle
changes, whether in terms of nutrition or
physical activity.

Some study limitations must be acknowl-
edged. First, blinding patients was not possible
in this study, which might have resulted in an
overestimation of the efficacy of the telemoni-
toring programme, as observed in other non-
blind studies [38]. Moreover, the impact of
being monitored according to the anticipated
study design is an unavoidable part of the pro-
cess and we cannot rule out the fact that patient
behaviour may have been different in real-life
settings. In addition, as in other randomised
trials, the results might not be generalised to
patients who would not have agreed to participa-
tion given their clinical or socio-economic profile (more serious medical history, poorer
health, limited internet access, etc.). Finally,
measuring satisfaction with a non-validated
questionnaire might have led to less reliable
measurements, making it more difficult to
compare the results to other studies.

To date, the long-term impact of telemoni-
toring has not been assessed to any considerable
extent in diabetes and, more generally, in
chronic diseases. It would therefore be inter-
esting to determine whether the positive trends
reported in our 1-year study are maintained in
the months following the cessation of tele-
monitoring. Further randomised studies should
be carried out to establish optimal modalities
for using telemonitoring devices in diabetes
(focusing on subject profiles, ideal time for
intervention, duration, combination or separa-
tion of telemonitoring and tele-education
devices, involvement of specialist doctors and
general practitioners, delegation of tasks to

Table 2 HbA1c changes according to randomisation group (N = 256) and frequency of device use

| Models | Over 1 year |
|--------|-------------|
|        | β   | 95% CI | p   | Δ   | 95% CI |
| Randomisation group | | | | | |
| Control group (CG) | | | | | |
| Telemonitoring group (TMG) | -0.16 | -0.32; 0.01 | 0.06 | * | -0.24 | -0.42; -0.07 |
| Number of connections (synthesis) | | | | | |
| Control group (CG) | ref | | | | |
| TMGs number ≤ median | -0.08 | -0.28; 0.13 | 0.45 | | -0.16 | -0.38; 0.05 |
| TMGs number > median | -0.23 | -0.43; -0.03 | 0.03 ** | | -0.31 | -0.53; -0.10 |
| Number of connections (Nutri-Educ) | | | | | |
| Control group | ref | | | | |
| TMGn number ≤ median | -0.11 | -0.31; 0.10 | 0.30 | | -0.19 | -0.40; 0.01 |
| TMGn number > median | -0.21 | -0.41; 0.00 | 0.05 ** | | -0.29 | -0.51; -0.07 |

Models adjusted for potential confounding factors defined a priori, taking into account time effect, centre effect and randomisation per stratum. HbA1c value at baseline was 7.75% (95% CI [7.65; 7.85])

With β linear regression coefficients, 95% CI 95% confidence intervals, p p value, m average HbA1c observed at T0 (%) and Δ predicted variation over 1 year

**Significant difference between groups (< 0.05); *difference at the limit of significance (< 0.10)
other healthcare professionals, etc.). Further improvement of the user profile definition should facilitate the targeting of individuals best suited to this type of tool and should allow personalisation of the intervention modalities.

CONCLUSION

This study shows that at-home telemonitoring and tele-education did not significantly improve glycaemic control in T2D subjects.

### Table 3: Changes in body weight and BMI by group and device use, stratified on gender

|                      | Weight (in kg) |       | 95% CI      | p   |       | 95% CI      | p   |
|----------------------|----------------|-------|-------------|-----|-------|-------------|-----|
|                      | β               |       |             |     | β     |             |     |
| In women (N = 86)    |                |       |             |     |       |             |     |
| Randomisation group  |                |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMG                  | −2.68          | −4.79 | −0.57       | 0.013** | −1.01 | −1.79; −0.23 | 0.011** |
| Connection (synthesis) |              |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMGs< number ≤ median| −1.84          | −4.46 | 0.79        | 0.170 | −0.75 | −1.72; 0.22  | 0.131 |
| TMGs> number > median| −3.39          | −5.87 | −0.91       | 0.007** | −1.23 | −2.16; −0.31 | 0.009** |
| Connections (Nutri-Educ) |            |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMGns< number ≤ median| −1.34          | −3.98 | 1.30        | 0.319 | −0.54 | −1.52; 0.44  | 0.282 |
| TMGns> number > median| −3.78          | −6.25 | −1.31       | 0.003** | −1.40 | −2.31; −0.48 | 0.003** |
| In men (N = 150)     |                |       |             |     |       |             |     |
| Randomisation group  |                |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMG                  | 0.46           | −0.87 | 1.80        | 0.498 | −0.16 | −0.27; 0.60  | 0.463 |
| Connection (synthesis) |              |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMGs< number ≤ median| 0.99           | −0.64 | 2.62        | 0.233 | −0.34 | −0.19; 0.88  | 0.208 |
| TMGs> number > median| −0.06          | −1.68 | 1.56        | 0.945 | −0.01 | −0.54; 0.52  | 0.966 |
| Connections (Nutri-Educ) |            |       |             |     |       |             |     |
| CG ref               | ref            |       |             |     | ref   |             |     |
| TMGns< number ≤ median| 0.91           | −0.70 | 2.52        | 0.268 | −0.32 | −0.21; 0.85  | 0.234 |
| TMGns> number > median| −0.01          | −1.65 | 1.63        | 0.987 | −0.001| −0.54; 0.54  | 0.996 |

Models adjusted for potential confounding factors defined a priori, taking into account time effect, centre effect and randomisation per stratum

*With β linear regression coefficients, 95% CI 95% confidence intervals, p p value

**Significant difference between groups (<0.05), * difference at the limit of significance (<0.10)
Despite a slight, albeit significant decrease in HbA1c levels in frequent device users as well as significant weight loss in women only. This strategy could contribute to new organisational modalities for diabetes management, making it possible to respond to current public health issues while maintaining the relevance and quality of follow-up. By offering an alternative to the conventional healthcare system, telemonitoring should allow the frequency of face-to-face medical consultations to be modulated according to individual requirements and availability, thus reducing healthcare costs. An economic analysis of the EDUC@DOM study after 2 years of follow-up will address the cost-saving impact of this strategy.

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**Fig. 3** Correlations between frequency of use and changes in HbA1c and BMI. With: solid lines for non-parametric (Lowess) models and dashed lines for linear models. Since the correlations are graphically linear, the following are presented: $\rho$, Pearson correlation coefficient, with confidence intervals estimated by bootstrapping (1000 replications) and $p$ value by the Pearson correlation coefficient test.

**Fig. 4** Evaluation of device use by telemonitored patients.
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**Data Availability.** The datasets generated and/or analysed during the current study are available from the corresponding author on reasonable request.

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