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Effects of the Proactive Interdisciplinary Self-Management (PRISMA) Program on Online Care Platform Usage in Patients with Type 2 Diabetes in Primary Care: A Randomized Controlled Trial

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Online care platforms can support patients with type 2 diabetes (T2DM) in managing their health. However, in the use of eHealth, a low participation rate is common. The Proactive Interdisciplinary Self-Management (PRISMA) program, aimed at improving patients’ self-management skills, was expected to encourage patients to manage their disease through the use of an online platform. Therefore, the objective of the current study was to investigate whether a group education program can improve the use of an online care platform in patients with T2DM treated by primary care providers in the Netherlands. In a randomized controlled trial, patients with T2DM received either PRISMA with usual care or usual care only. During a six-month follow-up period in 2014-2015, usage (number of log-ons and time spent per session) of an online care platform (e-Vita) aimed at improving T2DM self-management was assessed. A training about the functionalities of e-Vita was offered. The sample consisted of 203 patients. No differences were found between the intervention and control groups in the number of patients who attended the platform training (interested patients) ($X^2(1) = 0.58; p = 0.45$), and the number of patients who logged on at least once (platform users) ($X^2(1) = 0.46; p = 0.50$). In addition, no differences were found between the groups in the type of users—patients who logged on twice or more (active users) or patients who logged on once (nonactive users) ($X^2(1) = 0.56; p = 0.45$). The PRISMA program did not change platform usage in patients with T2DM. In addition, only a small proportion of the patients logged on twice or more. Patients probably need other encouragements to manage their condition using an online platform.
1. Introduction

Worldwide, the prevalence of diabetes mellitus is increasing dramatically. In the Netherlands, 66 per 1,000 persons have type 2 diabetes mellitus (T2DM), and this rate is expected to increase to 80 per 1,000 persons by 2025 [1]. Although patients with T2DM are primarily treated by primary care providers, this projected growth is expected to exceed the number of available providers [2]. Already, diabetes care providers see an increase in patients, which results in a decrease in face-to-face time available per patient. This has partly been tackled by a transfer of tasks, previously performed exclusively by general practitioners (GPs), to other medical professionals including specialized practice nurses (PNs). To deal with the increasing number of patients with T2DM and the burden of diabetes on healthcare, increased patient participation is needed, including more self-management. Self-management includes the active participation of patients in their treatment [3] to minimize the impact of chronic disease on their physical health and functioning and to enable patients to cope with the psychological effects of the illness [4]. Patient participation could be enhanced by offering them the possibility to track their own medical data together with tailored advice through eHealth [5].

eHealth applications, and more specifically online care platforms, provide the opportunity for self-management support and maintaining and/or improving the quality of chronic disease management by engaging patients in their own healthcare [6]. In general, online care platforms are environments in which patients can get an overview of their health outcomes, communicate with their care provider, and/or read information regarding their disease. It has been shown that such platforms are beneficial for people with T2DM [7]. These platforms have the potential to support patients in managing their own health and changing their lifestyle [8].

So far, the effects of online care platforms reported in systematic reviews vary [9–11]. Health behaviors and health-related outcomes have been shown to improve through the use of eHealth [12–15]. Moreover, these platforms were shown to be specifically beneficial for people with T2DM [7]. Therefore, platforms aimed at empowering patients can potentially decrease the workload of diabetes care providers and improve the (cost-)effectiveness of diabetes treatment [16, 17]. Nevertheless, implementation problems, nonadherence, and low participation are common [18–23]. A recent literature review of studies reporting online care platform use by patients with diabetes (type 1 and 2) revealed that 29% to 46% of them registered for a platform account; of those registered, 27% to 76% of patients used the platform at least once [23]. Platform use was associated with the following factors: patient characteristics (e.g., sociodemographic, clinical characteristics, and health literacy), technology (e.g., functionality, usability), and provider engagement. For facilitating the use of self-management support through a platform, patients first need to develop an intention for behavioral change, which can only be achieved if they have sufficient risk awareness, experience a need for behavioral change, and feel confident in making these changes [24]. In the Netherlands, an online care platform called e-Vita has been developed to improve patients’ self-management skills [25].

Group education could be a helpful way for patients to obtain an intention for behavioral change [26]. A recent systematic review suggests that group-based diabetes self-management education is associated with improved clinical and psychosocial outcomes [27]. The Proactive Interdisciplinary Self-Management (PRISMA) program is aimed at improving self-management skills in patients with T2DM [26]. PRISMA appeared to improve self-management behavior in terms of dietary behaviors, foot care, action planning, and medication adherence [24, 28]. In addition, a pilot study showed that the PRISMA program is promising for decreasing dietary intake in newly diagnosed, overweight patients with T2DM in secondary care [26]. PRISMA helps patients to evaluate their own risk factors, to set personal goals, and to formulate a realistic action plan. Therefore, the PRISMA program is expected to increase the patients’ motivation to behavior change and to motivate patients to manage their condition using an online platform.

The objective of the current study was to investigate whether a group education program aimed at empowering and stimulating self-management in patients with T2DM can improve the use of an online care platform.

2. Material and Methods

2.1. Study Design. The current study is part of the Diabetes Education and Self-Management to Increase Empowerment (DESTINE) study described in detail elsewhere [29]. DESTINE is a randomized controlled trial that followed enrolled patients for six months (Figure 1). The patients with T2DM received either the PRISMA program with usual care or usual care only. According to the guidelines of the Dutch College of GPs (NHG-Standard), usual care involves two to four visits per year with a PN and one annual check-up with a GP. All patients had access to the online care platform.

In another substudy of DESTINE, the effects of PRISMA on medication adherence were described [28]. Therefore, people 18 years old or older who diagnosed with T2DM and treated in primary care were included. Eight general practices in the eastern part of the Netherlands participated, and eligible patients were selected by GPs. Nonstratified block randomization was used to allocate participants to one of the two groups [30]. For the current study, we followed the methods of du Pon et al. [28].

2.2. Description of the Platform. In a previous study, the online care platform used for this study (e-Vita) showed that users of the platform had lower glycated hemoglobin (HbA1c) levels and reported higher quality of life, better well-being, lower diabetes-related distress, and better medication adherence than non-users [5]. However, the usage of this platform has been shown to be minimal in the past [5, 31]. Only 11% of the patients who were interested in the use of a care platform logged on at least once [31]. An improved version of the online platform e-Vita was used in our study. First, platform adjustments were conducted to
make the platform more user-friendly. Second, the log-on procedure was simplified. Third, a function regarding communication between patients and their PN was added. Finally, a training was offered to patients, their companions, and caregivers about the functionalities of e-Vita, as suggested by Roelofsen et al. [5].

The use of e-Vita was offered as part of a larger program that aimed to study the effects of an online platform for various chronic illnesses (T2DM, chronic obstructive pulmonary disease, and chronic heart failure). The platform (accessible through app.e-vita.nl) not only contains information about the patient’s health status but also offers participants more engaging options, such as formulating personal goals, participating in educational modules, exchanging messages between patient and care provider, or searching for information in the “library.” The language of the platform is Dutch. Table 1 shows these items in more detail.

In both groups, participants (and their spouses) had the option to use the platform. Therefore, participants were registered on e-Vita and were invited for a 90-minute training about this platform. In groups of 5–10, patients were introduced to e-Vita and received log-on data. Spouses were also invited due to their important role in supporting the patient who wish to access and use the platform [32]. They also became familiarized with the content by completing several exercises on the platform (e.g., “enter your weight” or “formulate a health goal”). After this training, participants were able to start using the platform immediately. The PNs also received training about how to use e-Vita to be able to respond to their patients’ messages, answer questions about the platform, and follow their patients’ activity in the educational modules.

2.3. Description of the Intervention. The intervention consisted of two group meetings about T2DM guided by care providers. The PRISMA program has been described in detail in du Pon et al. [29]. During PRISMA, patients were encouraged to set personal goals and formulate a realistic action plan. They could enter these personal goals on e-Vita and pursue their goals using the platform.
2.4. Outcomes. The outcome of this study was the change between groups in the usage of the e-Vita platform (number of log-ons and time spent per session), as assessed by user log files.

2.4.1. Usage of the e-Vita Platform. Each action performed by the users was logged in a file that was saved to the server and was available for the researcher. Log-in and log-out times of each session of all platform users were registered. First, for all platform users, the number of log-ons, the time spent per session, and the total time spent on the platform were calculated. Then, patients were defined according to what type of platform user they were (Figure 2) [25]. A session included all log-ons to the platform within 30 minutes [33].

2.4.2. Baseline Demographic and Clinical Data. Demographic baseline data were obtained from questionnaires, including sex, age, education level, health-related quality of life, emotional well-being, quality of received care, and eHealth literacy. Participants of the intervention group received the paper-based questionnaires immediately after the PRISMA program, whereas those of the control group received them by post.

Health-related quality of life was assessed by the EuroQol Five Dimension (EQ-5D-3L) scale [34]. This questionnaire consists of two parts: the EQ-5D descriptive system and the EQ visual analog scale (EQ-VAS). The EQ-5D-3L comprises five dimensions: mobility, self-care, usual activities, pain/discomfort, and anxiety/depression. Each dimension has three levels: no problems, some problems, and extreme problems. The EQ-5D-3L includes a visual analog scale (EQ-VAS) from which an individual rates their own health today from 0 (worst imaginable health) to 100 (best imaginable health). Emotional well-being was assessed by the 5-Item World Health Organization Well-Being Index (WHO-5) scale [35]. The WHO-5 covers five items: subjective quality of life based on positive mood (good spirits, relaxation), vitality (being active, waking up fresh and rested), and general interest (being interested in things). Each of the five items is rated on a 6-point Likert scale from 0 (not present) to 5 (constantly present). Higher scores represent higher levels of emotional well-being. Quality of received care was assessed by the HowRwe [36]. This questionnaire has four items concerning promptness, communication, personal relationship, and general satisfaction. Each item is scored using four levels ranging from "excellent" to "poor," and each level is assigned a score on a scale from 0 (poor) to 3 (excellent), with higher scores indicating a better patient experience.

eHealth literacy was assessed by the eHEALS questionnaire, an 8-item scale measuring perceived skills at finding, evaluating, and applying electronic health information to health problems [37]. For this study, the four most relevant items were used ($\alpha = .90$): (1) "I know how to find helpful health resources on the Internet"; (2) "I know how to use the Internet to answer my health questions"; (3) "I know what health resources are available on the Internet", and (4) "I feel confident in using information from the Internet to make health decisions". The items were measured with a 5-point Likert scale with response...
options ranging from “strongly disagree” to “strongly agree.” Total scores of the eHEALS were summed (possible range, 4–20), with higher scores representing higher self-perceived eHealth literacy.

In addition, baseline clinical data (T2DM duration, HbA1c, body mass index, and systolic blood pressure) were obtained from the personal health record systems of the GPs. Only data gathered less than four months before the intervention were used.

2.5. Sample Size Calculation. To show a difference of at least 20% in active users (15% in the control group versus 35% in the intervention group) for the primary outcome measure (usage of the e-Vita platform) with a two-sided risk alpha of 5% and a power of 80%, 81 individuals per group were needed using the unpoled Z-test. With an expected drop-out rate of 20%, we aimed to include 200 patients in our randomized controlled trial.

2.6. Analysis. Log files were collected for each patient over a six-month period. An intention to treat (ITT) analysis and a per protocol (PP) analysis were conducted. The PP analysis consisted of all patients of the intervention group who attended at least one session of the PRISMA program.

2.7. Statistical Analysis. All analyses were conducted using IBM SPSS Statistics version 22. Normally distributed data were presented as the means and standard deviation, whereas skewed data were presented as the medians and interquartile range. Dichotomous/categorical data were presented as numbers and percentage of the total. To evaluate differences in target variables (use of the online platform: number of log-ons and time spent per session) over time and between arms, the chi-square test and median difference scores (95% confidence intervals [CI]) were used. A subgroup analysis was conducted to investigate differences in platform use between male and female patients, younger (<65 years) and older patients (>65 years), and patients with a low eHEALS sum score (12 or less) and patients with a high eHEALS sum score (13 or more).

2.8. Ethics. This study was reviewed by the Medical Ethics Committee of Isala, Zwolle, the Netherlands. The committee decided that formal approval was not necessary (METC no. 14.07104).

3. Results

The inclusion lasted nine months (June 2014 to February 2015). Of 1,476 eligible patients, 203 (13.8%) were included in the study and signed the informed consent form; 101 patients were randomized to the intervention group and 102 were randomized to the control group. After inclusion, 10 patients (4.9%) withdrew from the study: 6 in the intervention group and 4 in the control group. Patients withdrew because of illness, immigration, or personal reasons. In the intervention group, 68 (71.6%) of 95 patients attended at least one of the two PRISMA meetings. The CONSORT patient flow chart is presented in Figure 3 [38].

3.1. Patient Characteristics. The patient characteristics are presented in Table 2. Of the total sample (n = 193), 60.1% were men. The mean age was 69.9 years (SD, 9.1; range, 35–96). No differences in patient characteristics were found between groups at baseline.

3.2. Platform Usage. The results of the ITT analyses on the platform usage are presented in Table 3.

Of the 193 participants in total, 58 (61.1%) patients of the intervention group and 65 (66.3%) patients of the control group registered on the platform during the training (interested patients). At 6 months, 33 (34.7%) patients of the intervention group and 33 (33.7%) patients of the control group had logged on to e-Vita once or more (platform users). Of the intervention group, 12 (12.6%) patients logged on once (nonactive users); 15 (15.3%) patients of the control group logged on once. Of the intervention group, 21 (22.1%) patients logged on twice or more (active users); 18 (18.4%) patients of the control group logged on twice or more. No differences were found between the groups in the number of interested patients ($X^2(1)=0.58; p=0.45$), the number of platform users ($X^2(1)=0.46; p=0.50$), or the type of users ($X^2(1)=0.56; p=0.45$).

Males more often were interested patient compared to females ($X^2(1)=7.70; p=0.00$). However, no association was found between sex and being a platform user ($X^2(1)=1.41; p=0.24$) or being an active user of the platform ($X^2(1)=0.83; p=0.36$). Also, no association was found between age and being an interested patient ($X^2(1)=0.12; p=0.73$), a platform user ($X^2(1)=0.26; p=0.61$), or active user of the platform ($X^2(1)=0.10; p=0.92$). In addition, no association was found between self-perceived eHealth literacy and being an interested patient ($X^2(1)=0.71; p=0.39$) or active user of the platform ($X^2(1)=0.62; p=0.43$). However, patients with a higher self-perceived eHealth literacy more often were platform users than patients with a lower self-perceived eHealth literacy ($X^2(1)=5.01; p=0.03$).

Of all patients, the median number of log-ons was 0 (interquartile range, 0–1) for the intervention group and 0 (interquartile range, 0–1) for the control group. Of the interested patients, the median number of log-ons was 2 (interquartile range, 1–7) for the intervention group and 2 (interquartile range, 1–5) for the control group with a median difference score of −0.34 (95% CI, −3.28 to 2.61). The total time spent on the platform was 56 minutes (interquartile range, 00:27–02:15) for the intervention group and 53 minutes (interquartile range, 00:27–02:14) for the control group with a median difference score of 0.90 (95% CI, −50.25 to 52.05).

The results of the PP analysis do not differ from the ITT results above (data not shown).

4. Discussion

The PRISMA group program, which aimed to improve T2DM patients’ self-management skills, was expected to increase patient motivation for behavior change and
encourage them to manage their disease through the use of an online platform. However, PRISMA did not result in a higher platform usage of patients with T2DM over a six-month period. No differences were found between the groups regarding the number of log-ons, the length of time spent per session, and the total time spent on the platform. In addition, attending the PRISMA intervention did not increase the chance of becoming a user of the platform or enhanced patient activity on the platform. Clearly, consideration of their own personal risk factors and choice of a specific behavior change goal during PRISMA did not encourage patients (enough) to actually use the platform.

Previous results on the effect of e-Vita usage on clinical outcomes were promising [25]. However, our study showed that usage of this platform was minimal. Even offering the PRISMA program coupled with a training to introduce the platform did not remove the threshold for patients to use the platform. Most of the patients who were registered for platform use and attended the training never logged on. Going online indeed requires extra effort for which the added value in view of self-managing a chronic disease using eHealth may not be automatically apparent for patients [39]. An explanation could be either a lack of worry about their future health [40, 41], resulting in an insufficient intrinsic motivation, and no intention to change behaviors. In addition, the absence of disease burden in early-stage T2DM could lead to a lack of patient motivation to use an online health platform. Most complications of T2DM arise on longer term. It would be easier to motivate patients if they could already see results of their self-management behavior in the short term. In addition, the mean age of the patient group was 70 years, and we did not find an association between age and platform use; however, according to another study, younger age is associated with platform use [23]. Although the Netherlands is a country with high levels of general Internet diffusion even among older adults [42], these new techniques may be not suitable for older generations who have not mastered the required computer skills [43]. We showed that male patients more often were interested in platform use compared to female patients, which corresponds to the findings of Roelofsen et al. [5]. Our finding that high eHealth literate patients used the platform more often than low eHealth literate patients corresponds to the literature [44, 45]. The improvement of eHealth literacy in
patients with T2DM needs attention. A recent literature review revealed the lack of data accuracy as the most important barrier to using eHealth for patients with T2DM [45]. This was often a result of manually reporting or the input of monitoring data. In our study, we recognized this problem. Patients probably need other encouragement to manage their T2DM through a platform, for example by follow-up meet-

ings, counsel sessions, or reminders. Otherwise, a platform might not be the most appropriate solution to improve self-management skills in patients with T2DM. The day-to-day management of T2DM can be complex and challenging and requires major responsibility of patients. The use of an online platform would be an extra activity and, as a result, outcomes of patient annual checkups were not displayed. Furthermore, despite our eHealth literacy. It is important to understand the importance of stimulating self-management skills in patients with chronic conditions via a platform. Other possible causes for the increase in platform usage could be the visual and technical improvements of the platform. In addition, bended care, where digital health and usual care are integrated, is likely to lead to increased use of an online program for patients with COPD [48]. Patients with diabetes log on to platforms for various reasons. According to the literature, frequently used features are viewing laboratory results [44], sending and reading messages [49], and participating in educational modules [50]; setting personal goals is the least popular feature [50].

4.1. Strengths and Limitations. To our knowledge, this is the first study that focuses on the extent to which patients use an online diabetes care platform after attending a group education program. The training offered about the functionalities of e-Vita helped patients to start their first session on the platform. In addition, communication between healthcare providers and patients, a new function on e-Vita, may have encouraged the use of the platform. Another strength of the study was that the participants were randomized over all general practices. This prevented influences from the general practices on the results. In addition, the study has pragmatic character, which means that it was designed to test PRISMA in the full spectrum of everyday usual care to maximize applicability and generalizability. Therefore, the study determined whether PRISMA actually increased the platform usage in real life. In 2017, the platform was renewed, extended with an app, and used by twenty care groups in the Netherlands.

Some limitations of this study need to be mentioned as well. The eight participating general practices diverged by inclusion rate: of the 1,476 invited patients, most were registered in three general practices. In addition, despite the instructions given during the training, the steps to gain access to the platform were considered complex. These factors posed a considerable barrier to using the platform, especially for those who were less able to handle computers. In the beginning of the project, technical problems appeared on the platform and, as a result, outcomes of patient annual checkups were not displayed. Furthermore, despite our efforts to enthuse patients about the PRISMA program, only 12% of the approached patients participated. Our target group might not be interested in this type of intervention, which could explain this low participation rate. In addition, most participants in our study were moderately or highly educated and reported a moderate or high eHealth literacy. Lower educated patients, in whom T2DM is most common [51], may not be interested in interventions such as PRISMA and e-Vita because of its complexity. Despite the randomized design, whether a patient becomes a (active) user of e-Vita could thus depend on caregiver attitudes toward participation, online platforms, and eHealth in general (selection bias).

5. Conclusions

The PRISMA program did result in a higher online healthcare platform usage in patients with T2DM, and also, the continuity of use was low. The added value of self-managing a chronic disease may not be automatically apparent for patients. In addition, platforms may not be suitable
for the older adults with lower eHealth literacy. There is a need to invest in developing online skills, particularly for older adults with lower eHealth literacy. Otherwise, these patients probably need other encouragements to help them manage T2DM using an online platform. Future research should explore other sources for patients to develop intentions to behavior change for facilitating the use of self-management support programs within a platform. Further development of the platform is ongoing.

Data Availability

The datasets generated and/or analyzed during the current study are not publicly available because public access was not included in the informed consent.

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper.

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Table 3: Results of platform usage by interest, user category, and activity.

| n (%)/mean ± SD/median (25–75 quartiles) | Intervention group (n = 95) | Control group (n = 98) | p value/median difference score (95% CI) |
|----------------------------------------|---------------------------|-----------------------|---------------------------------------|
| Interested patient^a                    | 58 (61.1)                 | 65 (66.3)             | 0.45^a                                |
| Platform user^b                         | 33 (34.7)                 | 33 (33.7)             | 0.50^b                                |
| Active user^c                           | 21 (22.1)                 | 18 (18.4)             | 0.45^#                                |
| Number of log-ons (all patients)        | 0 (0–1)                   | 0 (0–1)               | 0.77^#                                |
| Number of log-ons (only interested patients) | 2 (1–7)                 | 2 (1–5)               | 0.74^#                                |
| Range                                   | 1–11                      | 1–26                  | −0.34 (−3.28 to 2.61)                 |
| Categories                              |                           |                       | 0.85^*                                |
| 0–1 log-ons                             | 36 (63.2)                 | 46 (70.8)             |                                       |
| 2–4 log-ons                             | 9 (15.8)                  | 9 (13.8)              |                                       |
| More than 4 log-ons                     | 12 (21.1)                 | 10 (15.4)             |                                       |
| Time spent per session (hh/mm)          | 00 : 23 ± 00 : 09         | 00 : 25 ± 00 : 08     | 0.09^#                                |
| Total time spent (hh/mm)                | 00 : 56 (00 : 27–02 : 15) | 00 : 53 (00 : 27–02 : 14) | 0.93^#                                |
| Range                                   | 00 : 07–09 :02            | 00 : 06–09 :34        | 0.90 (–0.25 to 52.05)                 |

^*Pearson chi-square test. #Mann–Whitney U test. ^Atended the platform training. ^Logged on at least once after the training. ^Logged on for at least two sessions of 5 minutes or more after the training.

References

[1] N. Kleefstra, G. W. D. Landman, K. J. J. Van Hateren et al., “Dutch diabetes prevalence estimates (DUDE-1),” Journal of Diabetes, vol. 8, no. 6, pp. 863–865, 2016.
[2] World Health Organization, Global report on diabetes, World Health Organization, Geneva, 2016.
[3] T. Koch, P. Jenkin, and D. Kralik, “Chronic illness self-management: locating the “self”,” Journal of Advanced Nursing, vol. 48, no. 5, pp. 484–492, 2004.
[4] K. Lorig and H. Holman, “Arthritis self-management studies: a twelve-year review,” Health Education Quarterly, vol. 20, no. 1, pp. 17–28, 1993.
[5] Y. Roelofsen, S. H. Hendriks, F. Sieverink et al., “Differences between patients with type 2 diabetes mellitus interested and uninterested in the use of a patient platform (e-VitaDM-2/ZODIAC-41),” Journal of Diabetes Science and Technology, vol. 8, no. 2, pp. 230–237, 2014.
[6] P. M. Gee, D. A. Greenwood, D. A. Paterniti, D. Ward, and L. M. Soederberg Miller, “The eHealth enhanced chronic care model: a theory derivation approach,” Journal of Medical Internet Research, vol. 17, no. 4, article e86, 2015.
[7] M. Price, P. Bellwood, N. Kitson, I. Davies, J. Weber, and F. Lau, ”Conditions potentially sensitive to a personal health record (PHR) intervention, a systematic review,” BMC Medical Informatics and Decision Making, vol. 15, no. 1, article 159, p. 32, 2015.
[8] P. C. Tang, J. S. Ash, D. W. Bates, J. M. Overhage, and D. Z. Sands, ”Personal health records: definitions, benefits, and strategies for overcoming barriers to adoption,” Journal of the American Medical Informatics Association, vol. 13, no. 2, pp. 121–126, 2006.
[9] N. Van den Berg, M. Schumann, K. Kraft, and W. Hoffmann, ”Telemedicine and telecare for older patients—a systematic review,” Maturitas, vol. 73, no. 2, pp. 94–114, 2012.
[10] P. Sutcliffe, S. Martin, J. Sturt et al., ”Systematic review of communication technologies to promote access and engagement of
young people with diabetes into healthcare,” BMC Endocrine Disorders, vol. 11, no. 1, article 103, p. 1, 2011.

[11] S. J. Curry, W. Janssen, and M. A. Johansen, “Factors determining the success and failure of eHealth interventions: systematic review of the literature,” Journal of Medical Internet Research, vol. 20, no. 5, p. e10235, 2018.

[12] M. Mignerat, L. Lapointe, and I. Vedel, “Using telecare for diabetic patients: a mixed systematic review,” Health Policy and Technology, vol. 3, no. 2, pp. 90–112, 2014.

[13] M. S. Marcolino, J. X. Maia, M. B. M. Alkmim, E. Boersma, and A. L. Ribeiro, “Telemedicine application in the care of diabetes patients: systematic review and meta-analysis,” PLoS One, vol. 8, no. 11, article e79246, 2013.

[14] Z. Huang, H. Tao, Q. Meng, and L. Jing, “Management of endocrine disease. Effects of telecare intervention on glycemic control in type 2 diabetes: a systematic review and meta-analysis of randomized controlled trials,” European Journal of Endocrinology, vol. 172, no. 3, pp. R93–101, 2015.

[15] A. Steinsbekk, L. Ø. Rygg, M. Lisulo, M. B. Rise, and A. Fretheim, “Group based diabetes self-management education compared to routine treatment for people with type 2 diabetes mellitus. A systematic review with meta-analysis,” BMC Health Services Research, vol. 12, no. 1, article 2056, p. 213, 2012.

[16] K. Pal, S. V. Eastwood, S. Michie et al., “Computer-based diabetes self-management interventions for adults with type 2 diabetes mellitus,” Cochrane Database of Systematic Reviews, no. 3, article CD008776, 2013.

[17] A. Ramadas, K. F. Quek, C. K. Y. Chan, and B. Oldenburg, “Web-based interventions for the management of type 2 diabetes mellitus: a systematic review of recent evidence,” International Journal of Medical Informatics, vol. 80, no. 6, pp. 389–405, 2011.

[18] D. H. Gustafson and J. C. Wyatt, “Evaluation of ehealth systems and services,” BMJ, vol. 328, no. 7449, pp. 1150–1150, 2004.

[19] B. W. Hesse and B. Shneiderman, “eHealth research from the user’s perspective,” American Journal of Preventive Medicine, vol. 32, 5 suppl, pp. S97–S103, 2007.

[20] S. J. Curry, “eHealth Research and Healthcare Delivery: Beyond Intervention Effectiveness,” American Journal of Preventive Medicine, vol. 32, no. 5 suppl, pp. S127–S130, 2007.

[21] D. Flynn, P. Gregory, H. Makki, and M. Gabbay, “Expectations and experiences of eHealth in primary care: A qualitative practice-based investigation,” International Journal of Medical Informatics, vol. 78, no. 9, pp. 588–604, 2009.

[22] S. C. Wangberg, T. S. Bergmo, and P. J. M. Weijs, “Adherence in Internet-based interventions,” Patient Preference and Adherence, vol. 2, pp. 57–65, 2008.

[23] R. Sun, M. T. Korytkowski, S. M. Sereika, M. I. Saul, D. Li, and L. E. Burke, “Patient portal use in diabetes management: literature review,” JMIR Diabetes, vol. 3, no. 4, article e11199, 2018.

[24] M. Van Vught, M. de Wit, S. Bader, and F. J. Snoek, “Does low well-being modify the effects of PRISMA (Dutch DESMOND), a structured self-management-education program for people with type 2 diabetes?,” Primary Care Diabetes, vol. 10, no. 2, pp. 103–110, 2016.

[25] Y. Roelofs, S. H. Hendriks, F. Sieverink et al., “Design of the e-Vita diabetes mellitus study: effects and use of an interactive online care platform in patients with type 2 diabetes (e-VitaDM-1/ZODIAC-40),” BMC Endocrine Disorders, vol. 14, no. 1, article 233, 2014.

[26] A. J. Leibbrandt, J. C. Kiefte-de Jong, M. H. E. Hogendoelst, F. J. Snoek, and P. J. M. Weijs, “Effects of the PRo-active Interdisciplinary Self-MANagement (PRISMA, Dutch DESMOND) program on dietary intake in type 2 diabetes outpatients: a pilot study,” Clinical Nutrition, vol. 29, no. 2, pp. 199–205, 2010.

[27] A. Mohamed, E. Staite, K. Ismail, and K. Winkley, “A systematic review of diabetes self-management education interventions for type 2 diabetes mellitus in the Asian Western Pacific (AWP) region,” Nursing Open, vol. 6, no. 4, pp. 1424–1437, 2019.

[28] E. du Poon, S. El Azzati, A. van Dooren, N. Kleefstra, E. Heerdink, and S. van Dulmen, “Effects of a Proactive interdisciplinary self-management (PRISMA) program on medication adherence in patients with type 2 diabetes in primary care: a randomized controlled trial,” Diabetes Management, vol. 7, pp. 330–336, 2017.

[29] K. P. Suresh, “An overview of randomization techniques: an unbiased assessment of outcome in clinical research,” Journal of Human Reproductive Sciences, vol. 4, no. 1, pp. 8–11, 2011.

[30] Y. Roelofs, M. van Vught, S. H. Hendriks et al., “Demographical, clinical, and psychological characteristics of users and nonusers of an online platform for T2DM patients (e-VitaDM-3/ZODIAC-44),” Journal Diabetes Research, vol. 2016, article 634927, pp. 1–16, 2016.

[31] L. S. Mayberry, S. Kripalani, R. L. Rothman, and C. Y. Osborn, “Bridging the digital divide in diabetes: family support and implications for health literacy,” Diabetes Technology & Therapeutics, vol. 13, no. 10, pp. 1005–1012, 2011.

[32] F. Sieverink, S. Kelders, M. Poel, and L. van Gemert-Pijnen, “Opening the black box of electronic health collecting, analyzing, and interpreting log data,” JMIR Research Protocols, vol. 6, no. 8, article e156, 2017.

[33] M. Herdman, C. Gudex, A. Lloyd et al., “Measuring health utility using the WHO-Five well-being scale,” Quality of Life Research, vol. 15, no. 1, pp. 1005–1012, 2011.

[34] S. H. Hendriks, J. Rutgers, P. R. van Dijk et al., “Validation of the howRu and howRwe questionnaires at the individual patient level,” BMC Health Services Research, vol. 15, no. 1, 2015.

[35] C. D. Norman and H. A. Skinner, “eHEALS: the eHealth literacy scale,” Journal of Medical Internet Research, vol. 8, no. 4, article e27, 2006.
[39] T. Greenhalgh, S. Hinder, K. Stramer, T. Bratan, and J. Russell, "Adoption, non-adoption, and abandonment of a personal electronic health record: case study of HealthSpace," *BMJ*, vol. 341, no. nov16 1, p. c5814, 2010.

[40] M. Clark and S. E. Hampson, "Implementing a psychological intervention to improve lifestyle self-management in patients with type 2 diabetes," *Patient Education and Counseling*, vol. 42, no. 3, pp. 247–256, 2001.

[41] V. E. Coates and J. R. P. Boore, "The influence of psychological factors on the self-management of insulin-dependent diabetes mellitus," *Journal of Advanced Nursing*, vol. 27, no. 3, pp. 528–537, 1998.

[42] "CBS Statistics Netherlands," 2013, January 2019, https://opendata.cbs.nl/statline/#/CBS/nl/dataset/83429NED/table?fromstatweb.

[43] S. N. W. Vorrink, A. M. G. E. F. Antonietti, H. S. M. Kort, T. Troosters, P. Zanen, and J. J. Lammers, "Technology use by older adults in the Netherlands and its associations with demographics and health outcomes," *Assistive Technology*, vol. 29, no. 4, pp. 188–196, 2017.

[44] U. Sarkar, A. J. Karter, J. Y. Liu et al., "Social disparities in internet patient portal use in diabetes: evidence that the digital divide extends beyond access," *Journal of the American Medical Informatics Association*, vol. 18, no. 3, pp. 318–321, 2011.

[45] M. M. Alvarado, H. C. Kum, K. Gonzalez Coronado, M. J. Foster, P. Ortega, and M. A. Lawley, "Barriers to remote health interventions for type 2 diabetes: a systematic review and proposed classification scheme," *Journal of Medical Internet Research*, vol. 19, no. 2, article e28, 2017.

[46] E. Agardh, P. Allebeck, J. Hallqvist, T. Moradi, and A. Sidorchuk, "Type 2 diabetes incidence and socio-economic position: a systematic review and meta-analysis," *International Journal of Epidemiology*, vol. 40, no. 3, pp. 804–818, 2011.

[47] U. Öberg, U. Isaksson, L. Jutterström, C. J. Orre, and Å. Hörnsten, "Perceptions of persons with type 2 diabetes treated in Swedish primary health care: qualitative study on using eHealth services for self-management support," *JMIR Diabetes*, vol. 3, no. 1, article e7, 2018.

[48] E. P. Talboom-Kamp, N. A. Verdijk, M. J. Kasteleyn et al., "High level of integration in integrated disease management leads to higher usage in the e-Vita study: self-management of chronic obstructive pulmonary disease with web-based platforms in a parallel cohort design," *Journal of Medical Internet Research*, vol. 19, no. 5, article e185, 2017.

[49] M. Tenforde, A. Nowacki, A. Jain, and J. Hickner, “The association between personal health record use and diabetes quality measures,” *Journal of General Internal Medicine*, vol. 27, no. 4, pp. 420–424, 2012.

[50] F. Sieverink, S. M. Kelders, L. M. A. Braakman-Jansen, and J. W. C. van Gemert-Pijnen, “The added value of log file analyses of the use of a personal health record for patients with type 2 diabetes mellitus preliminary results,” *Journal of Diabetes Science and Technology*, vol. 8, no. 2, pp. 247–255, 2014.

[51] "National Institute for Public Health and the Environment [Dutch: Rijksinstituut voor Volksgezondheid en Milieu] (RIVM)," 2019, https://www.volksgezondheidenzorg.info/onderwerp/diabetes-mellitus/cijfers-context/huidige-situatie#node-diabetes-naar-opleiding(January.03.2019).
