The effect of motivational interviewing on glycaemic control and perceived competence of diabetes self-management in patients with type 1 and type 2 diabetes mellitus after attending a group education programme: a randomised controlled trial

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Abstract

Aims/hypothesis The aim of this study was to measure the efficacy of motivational interviewing (MI) compared with usual care on changes in glycaemic control and competence of diabetes self-management in patients with diabetes mellitus.

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Methods Patients were eligible if they had type 1 or 2 diabetes mellitus, were over 18 years of age and had participated in a 4 day group education programme offered at a diabetes clinic at a university hospital in Denmark. Exclusion criteria included pregnancy, severe debilitating disease and cognitive deficit. Out of 469 patients who attended the group education programme, 349 patients were randomised to either a usual care control group or an intervention group, which received up to five individual counselling sessions in 1 year based on MI, in addition to usual care. A randomised parallel design was used and open-label allocation was done by random permuted blocks, with allocation concealment by sequentially numbered, sealed, opaque envelopes. The primary outcome was glycated haemoglobin (HbA_{1c}). Analysis regarding measurements of glycated haemoglobin (HbA_{1c}) and competence of self-management (using the Problem Areas in Diabetes Scale [PAID] and Perceived Competence for Diabetes Scale [PCDS]) was based on 298 participants followed for a 24 month period. Data were collected at the Department of Endocrinology at Odense University Hospital. Our hypotheses were that MI could: (1) reduce HbA_{1c} levels; (2) increase self-efficacy; and (3) increase diabetes self-care, compared with usual care.

Results Out of the 176 included in the control group and 173 in the intervention group, 153 and 145 were analysed in the groups, respectively. When using the baseline value as covariate there were no significant differences in change score between the two study groups with regard to mean level of HbA_{1c} (0.131, $p=0.221$), PAID scores (-1.793 , $p=0.191$) or PCDS scores (0.017, $p=0.903$) at the 24 month follow-up, using a mixed effects regression model. The

patients in the intervention group showed significantly higher levels of perceived competence in dealing with diabetes compared with the control group (mean change score = -0.387 , $p=0.002$) following 1 year of intervention. *Conclusion/interpretation* We were unable to demonstrate any benefit, over or above usual care, of MI in patients with diabetes who have just completed a diabetes education programme, and who have well-regulated diabetes.

Trial registration Clinical Trials NCT00555854.

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Keywords Clinical trial · Diabetes mellitus · Motivation · Patient education · Self care

Abbreviations

MI	Motivational interviewing
MITI	Motivational Interviewing Treatment Integrity
OHA	Oral hypoglycaemic agents
PAID	Problem Areas in Diabetes
PCDS	Perceived Competence for Diabetes Scale

Introduction

Self-management of diabetes requires that patients are able to reconcile their resources and preferences with the therapeutic regimen of diabetes, which can be a challenging task for many patients [1, 2]. It has been widely accepted that diabetes education is not only required in the first few months following diagnosis but is an important component of ongoing diabetes care because of the onerous requirements for self-care that demand multiple daily decisions in order to balance diet, physical activity and medications [3]. There has been a keen interest in examining the impact of different kinds of patient education programmes on diabetes self-management. The findings of several meta-analyses of randomised controlled trials provide extensive evidence for the effectiveness of behavioural and educational intervention on fasting blood glucose and glycated haemoglobin, but the long-term effects of such interventions are uncertain [4–8]. Knowledge about the effectiveness of behavioural and educational intervention on other diabetes-related outcome measures, including blood pressure, lipid profile, body weight, self-management skills, health behaviour and psychosocial aspects, is less clear [3, 9, 10]. Critical assessment of the impact of behavioural and educational programmes requires further research based on rigorous methods in high quality studies with a large number of participants, long-term follow-up on effectiveness and well-defined interventions [9–11].

Motivational interviewing (MI), defined by Rollnick and Miller, is a well-defined and scientifically tested method of client counselling that has successfully been used to elicit and sustain a person's behaviour changes in a number of healthcare areas [12]. A recent review of MI showed improvements in health behaviour (e.g. diet and exercise) in patients with diabetes [13]. Research has not yet been able to establish full consensus on the effect of MI on clinical [14–19] and psychosocial aspects [15, 18, 20–22] in diabetes. Previous reviews emphasise the need for studies of high methodological quality and adequate power to explore the effect of MI on glycaemic control and well-being in patients with diabetes [11, 13, 23].

In this study, we explored the significance of supporting patients with diabetes to manage their diabetes over time. The aim was to study the effect of a 1-year intervention programme based on MI following a group education programme on glycaemic control and competence of management in patients diagnosed with type 1 or type 2 diabetes mellitus. Our hypotheses were that, in a sample of people who have just completed a group education programme, a course of motivational intervention following group education could: (1) reduce HbA_{1c} levels; (2) increase self efficacy; and (3) increase diabetes self-care, compared with usual care.

Method

Research settings

This study was conducted at a diabetes clinic, located within a university hospital in Denmark with an intake of patients from both urban and rural areas in the region. The hospital is a local hospital with an intake area that reflects other hospital intake areas in Denmark. Patients can be referred by their general practitioner or other hospital departments to the diabetes clinic. At the clinic, patients are offered ongoing individual counselling by dietitians, nurses or physicians and group education delivered by a multidisciplinary healthcare team over 4 days. The 4 day diabetes education programme includes approximately eight to ten patients and focuses on a range of themes such as diabetes treatment, prevention of diabetes-related complications, self-monitoring of blood glucose, diet, physical activity, alcohol use and issues regarding insurance. The teaching method is based on guided self-determination designed to guide patients and professionals in mutual problem-solving [24].

Research participants and design

Patients were eligible if they had type 1 or type 2 diabetes mellitus, were over 18 years of age and had participated in

the group education programme offered at the diabetes clinic. Exclusion criteria included pregnancy, severe debilitating disease and cognitive deficit. The patients were informed at day 1 and recruited at day 4 while participating in the group education programme. Following informed consent, the patients were randomised to either a control group or an intervention group. Randomisation was done immediately after participation in the group education programme. No stratification was used. All outcome measures were assessed at randomisation, 1 and 2 years after randomisation in both groups. Randomisation was generated by random permuted blocks, with allocation concealment by sequentially numbered, sealed, opaque envelopes. The person generating the allocation scheme did not administer the allocation of the patients to the two groups and was not part of the research team.

The sample size was determined by a power calculation. With 352 patients, 176 in each group, the trial can detect a 0.4% difference in HbA_{1c}. The power was set to 90%. This calculation was based on a standard deviation of 1.15 in the HbA_{1c} value and a 5% two-sided significance level.

Usual care

Medical treatment was not part of the intervention. All participants, irrespective of participation in the intervention group or the control group, therefore underwent the same routine check-up at their general practitioner or outpatient clinic in charge of their diabetes care. This usually involved four physician visits per year. Biochemical tests and examinations were usually performed during the visits in accordance with national diabetes guidelines. Individual counselling and recommendations based on the results of the examinations, biochemical tests and their self-monitoring of blood glucose was given. Renewal of prescribed medication and test strips for blood glucose monitoring were also given at these check-ups. Patients could be referred for individual counselling in change of diet, physical activity, smoking habits and alcohol abuse if required by their usual healthcare provider.

Research intervention

The theoretical approach of the intervention was based on self-efficacy theory and MI spirit. Perceived self-efficacy is defined as people's beliefs about their capabilities of producing designated levels of performance exercising influence over events that affect their lives [25]. MI was used as a method to facilitate this process. MI is a directive counselling style for eliciting behaviour change by helping patients to explore and resolve ambivalence [26, 27].

In addition to usual care, patients in the intervention group received a 1 year MI programme consisting of five

individual counselling sessions lasting approximately 45 min and offered at 1, 3, 6, 9 and 12 months. Each participant in the intervention group had a healthcare professional assigned to them who was trained in MI. The style of the interview was: (1) seeking to understand the person's frame of reference; (2) expressing acceptance and affirmation; (3) eliciting and selectively reinforcing the client's own self-motivational statements of problem recognition, concern, desire and intention to change, and ability to change; (4) exploring the client's degree of readiness to change; and (5) affirming the client's freedom of choice and self-direction. The role of exploring readiness to change was that this was used as a component of the therapeutic process and not an outcome. Each session followed a semi-structured interview format of MI, especially developed for this intervention programme. Participants brought up any problematic issues related to diabetes self-care during sessions. The participants in the intervention group could be referred by the healthcare professional to individual counselling in changes of diet, a smoking cessation programme, counselling in alcohol abuse and an exercise programme, as they required.

Education of the healthcare professionals prior to the intervention

Three diabetes specialist nurses, two dietitians, one physiotherapist and one psychologist were educated to carry out MI. They were all coached by a MI trainer from the Motivational Interviewing Network of Trainers in the Nordic countries. The course comprised a 5 day theoretical introduction to strategies in MI. The course was followed by three practical coaching sessions every 3 months for 18 months. The theoretical and practical part of the education included training in the key elements of MI, which is generally facilitating through eliciting change talk and exploring ambivalence about behavioural change, while trying to examine discrepancies between the individual's current behaviour and core values or personal goals. The healthcare professionals were introduced to the MI method including reflective listening and acknowledgement to allow them to be able to clarify the patient's goals and concerns and elicit reasons for change in the patient's own words. The role of the healthcare professionals was to coach and support the patient in discovering and developing his/her own resources for change and management at the patient's request.

After the 5-day course, the healthcare professionals were individually supervised by the MI trainer in ten real patient situations for 1 year. The supervision included videotaping and evaluation inspired by the Motivational Interviewing Treatment Integrity (MITI) coding system (T. B. Moyers, T. Martin, J. K. Manuel, W. R. Miller, D. Ernst, 2007;

available to trainers and members of the Motivational Interviewing Network of Trainers [MINT]). The MITI coding system is divided into a global rating and behavioural counts. The global rating is a five-point Likert scale, where 1 indicates low competence in MI and 5 indicates high competence in MI. The behaviour counts reveal MI behaviour in proportion to all behaviour, where a high percentage indicates a high competence in MI.

Measurements

Questionnaires Competence of diabetes self-management was assessed by validated questionnaires translated into a Danish version in accordance with recommendations [24]. The Problem Areas in Diabetes Scale (PAID) was used to measure diabetes-related distress reported by the participants. This 20-item questionnaire assesses a wide range of feelings related to living with diabetes and diabetes treatment [28, 29]. Each item can be rated on a five point Likert scale ranging from 0 (not a problem) to 4 ('a serious problem'). When transforming PAID scores into a 0–100 scale, all item scores are summed and multiplied by 1.25, which results in an overall PAID score with higher scores indicating greater emotional distress [30, 31]. High PAID scores are associated with low self-reported adherence [29]. The Perceived Competence for Diabetes Scale (PCDS) was used to measure competence at carrying out the diabetes treatment regime. This is a five item questionnaire assessing the degree to which participants feel confident about dealing with diabetes [32, 33]. Each item can be rated on a seven point Likert scale ranging from 1 ('not true at all') to 7 ('very true'). The score on the PCDS is calculated by averaging the responses on the five items. The minimum average score is the one equalling the lowest possible perceived competence in dealing with diabetes and the maximum average score is 7, equalling the highest possible perceived competence in dealing with diabetes.

Laboratory measurements One laboratory analysed all the blood samples. Glycaemic control was assessed using HbA_{1c}, which was measured by a high-performance liquid chromatography-based ion exchanged procedure (Tosho Alc 2.2, Tokyo, Japan). The reference range was 4.3–6.3%. Total cholesterol, HDL cholesterol and triacylglycerol levels were measured in serum by enzymatic methods (Boehringer Mannheim Diagnostica, Mannheim, Germany). LDL-cholesterol was calculated by Friedewald's equation [34].

Clinical measurements Both height and weight were measured without shoes. Weight was measured without accessories, i.e. wallet, keys, mobile phone. Waist circumference was measured with a measuring tape directly on the individual's skin. Restrictive garments and clothing were removed. The

tape was placed at the uppermost border of the iliac crest horizontal to the floor. Measurement was made at the end of a normal expiration. Blood pressure was measured by the auscultatory method with use of a stethoscope and a sphygmomanometer. An inflatable cuff was placed around the upper left arm, at the same vertical height as the heart. Measurement was made at rest in a sitting position.

Statistical analysis

For data analyses, SPSS (version 18, Chicago, IL, USA) for Windows and STATA (version 11, College Station, TX, USA) were used.

The baseline values are reported as means±SD or *n* (%). On the basis of an intention to treat analysis, a mixed effects regression model with random intercept was used to compare primary and secondary outcomes at 12 and 24 months between the intervention and control groups, where treatment was taken as the explanatory variable and the baseline value as a covariate. Further, each outcome was adjusted for the covariates, age, sex, type of diabetes, level of education, HbA_{1c} level at baseline, and healthcare professionals delivering the intervention. A random intercept was included in order to account for possible within-individual dependence due to repeated measurements. In the same fashion, a random intercept was included to account for cluster effects due to the healthcare professionals. Results are presented as means (95% CI). A *p* value of <0.05 was regarded as statistically significant.

Research ethics

The study design was approved by the Regional Scientific Ethics Committee of Southern Denmark (project-ID: VF-20050131) and the study was reported to the Danish Data Protection Agency. All patients were informed about the aim of the study and were included in the study after informed consent. The study was performed in accordance with the Declaration of Helsinki.

Results

Baseline data

In the period between December 2005 and June 2009, 469 patients attended the diabetes education programme at the diabetes clinic. Some 115 patients declined to participate in the trial and five patients did not meet the inclusion criteria. The reasons for declining were mainly lack of time or lack of energy to complete the trial. Of the 120 non-participants, 62 were women and 56 were men. The distribution of sex

was comparable in the non-participant group and the eligible group of patients. The mean age in the non-participant group was 59.1 ± 11.8 years. This was significantly higher ($p=0.040$) than the age of those who participated in the trial, where the total mean age was 56.4 ± 12.1 years. The duration of diabetes was 7.1 ± 10.0 years in the non-participant group and 4.7 ± 6.7 years in total among the participants. This difference was significant ($p=0.008$). No difference was found between the non-participant patients and the eligible patients with regard to weight, blood pressure, HbA_{1c} and lipid profile.

Of the 349 eligible patients, 173 patients were randomised to the intervention group and 176 to the control group. During the 2 year follow-up period, 51 participants (15%) dropped out, of whom 28 had been allocated to the intervention group and 23 to the control group (Fig. 1). The baseline characteristics and health status of both groups were similar (Table 1). We found no difference in the characteristics of dropout participants compared with those who remained in the study, except for the mean age, where the dropouts were younger than the intervention group (mean difference 4.50 years, $p=0.035$).

Intervention

In the intervention group, the average visits per patient were 4.6 with an average time of 34 min per session during the 1 year intervention programme. In the intervention group, 85% completed the full intervention programme

with the five sessions. The last 15% in the intervention group had between one and three sessions.

In the intervention group, 17% were referred to individual counselling in change of diet. The mean numbers of visit per patient were 2.0 (minimum one and maximum five visits). Some 25% in the intervention group were referred to an exercise programme. The mean numbers of visits per patient were 5.7 (minimum one and maximum 50 visits). Of the patients who participated in dietary counselling and the exercise programme, 15% received both therapies. Seven per cent in the intervention group participated in individual counselling on smoking cessation. The mean number of visits per patient was 4.9 (minimum 2 and maximum 10 visits). Finally, 3% of the patients in the intervention group were referred to individual counselling in change of alcohol use. No monitoring in the control group was done.

The global rating scores in the MITI coding of the videotapes were between 4 and 5 points. The behaviour counts showed 80–100% MI adherent behaviour in proportion to the total behaviour, 76–83% of open questions in proportion to all questions asked, 36–42% of complex reflections in proportion to all reflections made and the reflection to question ratio was between 0.8 and 1.8.

Random effects analysis

When using the baseline value as covariate there were no significant differences in change in score between the two

Fig. 1 Flow diagram of inclusion of the study population

Table 1 Characteristics of the patients at baseline

Variable	Intervention (<i>n</i> =173)	Control (<i>n</i> =176)
Sex		
Male	90 (52)	86 (49)
Female	83 (48)	90 (51)
Age (years)	57.1±12.6	55.8±11.6
Diabetes duration (years)	4.7±6.9	4.7±6.5
Type of diabetes		
Type 1	40 (23)	36 (21)
Type 2	133 (77)	140 (79)
Diagnosed retinopathy	25 (15)	20 (11)
Educational level ^a		
≤Middle school	122 (70)	120 (68)
≥High school	51 (30)	56 (32)
Connection to the labour market		
Transfer payment	109 (63)	106 (60)
Employed/self-employed	64 (37)	70 (40)
Civil status		
Live alone	55 (32)	47 (27)
Cohabiting	118 (68)	129 (73)
Weight (kg)	89±18	90±19
Body mass index (kg/m ²)	30.8±5.8	31.1±6.3
Waist circumference (cm)	102±15	102±15
HbA _{1c} (%)	7.0±1.2	7.0±1.2
Blood pressure (mmHg)		
Systolic	131.9±16.7	132.5±17.1
Diastolic	75.4±10.4	76.3±9.7
Cholesterol (mmol/l)		
Total	4.4±1.1	4.5±1.0
LDL	2.3±1.0	2.3±0.9
HDL	1.4±0.4	1.4±0.5
Triacylglycerol	1.6±1.1	1.9±1.4
PAID	20.0±17.7	19.6±16.3
PCDS	6.3±1.0	6.1±1.1
Hypoglycaemic treatment		
Insulin	47 (27)	52 (30)
OHA	80 (46)	74 (42)
Insulin and OHA	14 (8)	21 (12)
Antihypertensive treatment	104 (60)	109 (62)
Cholesterol-lowering treatment	107 (62)	119 (68)

Data are mean±SD or *n* (%)^a Middle school is between seventh and ninth grade. High school is ninth to twelfth grade

study groups with regard to mean level of HbA_{1c} (0.131, $p=0.221$), total cholesterol (0.154, $p=0.110$), LDL cholesterol (0.001, $p=0.981$), PAID scores (-1.793 , $p=0.191$) or PCDS scores (0.017, $p=0.903$) at the 24 month follow-up (see Table 2). Also, no statistically significant differences were found in weight (Δ mean change score=0.664, $p=0.291$), waist circumference (Δ mean change score=1.366, $p=0.297$), systolic blood pressure (Δ mean change score= -0.145 , $p=0.927$) or diastolic blood pressure (Δ mean change score= -0.006 , $p=0.995$) between the

intervention and control group at the 24 month follow-up (Table 2). The patients in the intervention group showed significantly higher levels of perceived competence in dealing with diabetes compared with the control group (Δ mean change score= -0.387 , $p=0.002$) 12 months post intervention.

We found no statistically significant effect between the healthcare professionals delivering the intervention.

Similar changes in medical treatment over time were found in the intervention and control group as regards insulin, oral hypoglycaemic agents (OHAs), antihyperten-

Table 2 Changes in primary and secondary outcome measures over 12 and 24 months in intervention group and control group

Measure	Unadjusted mean scores±SD Baseline	Adjusted mean scores (95% CI)		Analysis results ^b			
		12 months	24 months	ΔMean ^a 12 months	<i>p</i> _{12 months}	ΔMean ^a 24 months	<i>p</i> _{24 months}
HbA _{1c} (%)							
Intervention	7.02±1.20	6.93 (6.80–7.06)	7.16 (7.01–7.31)	0.029	0.757	0.131	0.221
Control	7.03±1.16	6.96 (6.83–7.08)	7.29 (7.15–7.44)				
Total cholesterol (mmol/l)							
Intervention	4.43±1.08	4.27 (4.16–4.39)	4.37 (4.24–4.51)	0.073	0.375	0.154	0.110
Control	4.54±1.01	4.41 (4.30–4.52)	4.59 (4.46–4.72)				
LDL-cholesterol (mmol/l)							
Intervention	2.35±0.99	2.31 (2.21–2.40)	2.43 (2.32–2.54)	0.008	0.908	0.001	0.981
Control	2.30±0.86	2.32 (2.32–2.42)	2.44 (2.33–2.55)				
PAID							
Intervention	20.0±17.7	18.5 (16.3–20.6)	19.5 (17.6–21.4)	1.522	0.321	−1.793	0.191
Control	19.6±16.3	20.2 (18.1–22.3)	17.9 (16.1–19.8)				
PCDS							
Intervention	6.27±1.05	6.25 (6.07–6.42)	6.03 (5.84–6.22)	−0.387	0.002	0.017	0.903
Control	6.18±1.11	5.82 (5.65–5.98)	6.00 (5.81–6.19)				
Weight (kg)							
Intervention	89.7±17.7	89.3 (88.5–90.1)	90.8 (89.9–91.6)	0.588	0.210	0.664	0.291
Control	89.6±18.3	90.3 (89.5–91.0)	91.8 (90.9–92.6)				
Waist circumference (cm)							
Intervention	102.2±14.6	102.8 (102.0–103.6)	102.4 (100.6–104.3)	0.230	0.690	1.366	0.297
Control	102.4±15.1	102.9 (102.1–103.7)	103.7 (101.9–105.5)				
Systolic BP (mmHg)							
Intervention	131.9±16.7	135.0 (132.6–136.9)	133.3 (131.1–135.5)	−2.33	0.102	−0.145	0.927
Control	132.5±17.1	132.6 (130.7–134.6)	133.1 (131.0–135.2)				
Diastolic BP (mmHg)							
Intervention	75.4±10.5	78.9 (77.6–80.2)	77.1 (75.8–78.4)	−1.040	0.260	−0.006	0.995
Control	76.3±9.7	78.1 (76.9–79.4)	77.3 (76.1–78.6)				

^a The difference in mean change score between groups ($[\text{Int}_0 - \text{Int}_1] - [\text{Con}_0 - \text{Con}_1]$)

^b Comparison between the two groups for primary and secondary outcomes was performed by a mixed effects regression model with random intercept, adjusting for baseline values and age, sex, type of diabetes, level of education and HbA_{1c} level at baseline with a random intercept

sive agents, cholesterol-lowering agents, anticoagulants and steroids (electronic supplementary material [ESM] Table 1).

Discussion

The main findings of this 1 year study were that patients with diabetes participating in a MI programme did not improve glycaemic control or competence of self-management in the intervention group compared with the control group. We found that the intervention maintained patients' perceived competence in dealing with diabetes compared with the control group just after the intervention. However, this difference between the groups disappeared at the 24 month follow-up.

The lack of effect on HbA_{1c}, lipid profile and other clinical outcomes such as weight and waist circumference might indicate that the patients in the intervention group did not change behaviour towards a higher degree of diabetes self-care including healthy diet and more physical activity. These findings are supported by the fact that we found no evidence that MI improved competence of self-management over time. This might indicate that it takes more time before the effectiveness of MI on patients' cognition could be seen in desirable behavioural changes in diabetes self-management [35].

Previous meta-analyses have revealed that behavioural and educational interventions in diabetes have produced modest short-term improvements in glycaemic control. There was a tendency to non-significant improvements in

glycated haemoglobin in the included studies where the mean HbA_{1c} values were less than 8.5% at baseline [4–6, 36]. A tendency to a reduced effect on glycaemic control was found in studies where the duration of diabetes was above 9 months [6]. Our results correspond with these findings. Essentially, MI might not improve glycaemic control in patients who have a relatively lower HbA_{1c} and longer duration of diabetes, especially when general practitioners or diabetologists see them regularly.

It appears from the mean PAID result at baseline that patients entering into the study had a relatively low level of diabetes-related distress. This is similar to what is found in unselected outpatient groups [31]. The fact that the patients had participated in an education programme before entering into the study might be the reason for such a relatively low level of diabetes-related distress at baseline, which might be reflected in the follow-up measurement. The mean PCDS results at baseline were close to the maximum score in both the intervention and the control group, which might also be explained by the fact that the patients had received education in diabetes before entering the study. Patients in the intervention group scored significantly better in perceived competence in dealing with diabetes just after the end of the intervention, compared with the control group. This might indicate that the intervention helped patients feel more competent in managing their diabetes. However, the instrument has not been developed directly to be used in clinical practice and the interpretation of how changes in mean PCDS score have an influence on patients' diabetes self-management is not clear. Moreover, the ceiling effect occurred as a high proportion of participants in the study had maximum scores in PCDS at baseline.

This study provided insight into the effect of a behavioural and educational intervention. Implementing MI as a new method in diabetes care required a shift in the healthcare professionals' role as well as practical training. Previous studies have shown that several barriers have to be overcome before a new method can be applied [37–39]. To accommodate these barriers, supervision and practical coaching sessions were held after the theoretical introduction to MI. It is a strength of this study that there has been supervision, which included videotaping and evaluation by the MITI coding system, both to facilitate the use of MI and to ensure that the method was used correctly in the intervention. The evaluation using MITI coding demonstrated a high competency in using MI among the healthcare providers in the study.

Limitations

The power calculation showed that 352 patients were needed to show a 0.4% difference in HbA_{1c}. Unfortunately, several patients declined participation, which resulted in

lower statistical power than planned. The fact that we failed to achieve our recruitment target might have hindered us statistically in detecting a possible intervention effect and thus we risked overlooking a real difference between intervention and control group. Furthermore, we cannot totally exclude the risk of contamination bias, which in this case could occur if patients in the control group changed their behaviours after interacting with patients in the intervention group, or if the healthcare professionals assigned to deliver the intervention also treated patients in the control group coming to the diabetes clinic for routine care. Contamination bias might diminish the magnitude of the difference between the intervention and usual care. Most of the patients entering this study, however, went to their general practitioner for routine care, which might minimise the risk of contamination bias. Nevertheless, use of a cluster design might have minimised the possibility of contamination bias even more.

Significance

With this randomised controlled trial we were unable to provide evidence that MI should be considered for adoption in routine diabetes care. When implementing a new approach in diabetes treatment, we have to consider who are most likely to benefit from this kind of intervention. Considering the group of patients who declined to participate in our trial, it was evident that they were older and had had diabetes for a longer time period than those who participated in the trial. This could indicate that intensive interventions might be less attractive and manageable for the older population with diabetes. We also might speculate that behavioural interventions including MI are only effective for problem behaviours if these are associated with poor glycaemic control.

In our trial, no improvements were found in cardiovascular risk profiles, whereas such improvements are often found in pharmacological studies [40, 41] with consequent improvements in mortality and morbidity. We did not intervene in medical treatment and it is therefore reasonable to assume that it was the intervention based on MI that improved the patient's perceived competence in diabetes post intervention. This might indicate that behavioural and educational intervention adds another dimension to diabetes care, compared with medical treatment. Therefore, the effectiveness of behavioural and educational intervention given in addition to usual care might be best assessed with psychological rather than biochemical variables in patients with HbA_{1c} levels less than 8%. This is substantiated by a previous review that pointed out the difficulty in assessing the unique contribution of diabetes patient education to diabetes outcomes [42]. Another review recommended that the evaluation of diabetes patient education, including behavioural and educational intervention, is concentrated on the goal of optimal adjustment to living with diabetes, as

this kind of treatment is believed to play a discernable role in knowledge, self-determination, self-management and psychological adjustment [43].

MI proved to be a good power-generating tool for the staff members, who need both knowledge and practical communication tools to talk to patients about difficult topics. Based on this knowledge, the professions involved in clinical practice can be advanced by a more qualified reinforcement of initiatives regarding behavioural changes concerning lifestyle issues in a clinical setting. To succeed with this, it is necessary to focus on the implementation process and continuously follow up on this to achieve this goal.

Finally, we were unable to demonstrate any benefit of MI over and above usual care in patients with well-regulated diabetes and newly completing a diabetes education programme.

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