

Cardiovascular Risk Education and Social Support (CaRESS): Report of a Randomized Controlled Trial from the Kentucky Ambulatory Network (KAN)

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Purpose: Test a practice-based intervention to foster involvement of a relative or friend for the reduction of cardiovascular risk in patients with type 2 diabetes.

Methods: We enrolled in a randomized controlled trial 199 patients and 108 support persons (SPs) from 18 practices within a practice-based research network. All patient participants had type 2 diabetes with suboptimal blood pressure control and were prepared to designate a SP. A subset of the patients also had dyslipidemia. All study visits were conducted at the practice sites where staff took standardized blood pressure measurements and collected blood samples. All patients completed one education session and received newsletters aimed at improving key health behaviors. Intervention group patients included their chosen SP in the education session and the SPs received newsletters.

Results: After 9 to 12 months, the intervention had no significant effect on systolic blood pressure, HbA1C, health-related quality of life, patient satisfaction, medication adherence, or perceived health competence. Power was insufficient to detect an effect on low-density lipoprotein cholesterol. Baseline cardiovascular risk values were not very high, with mean systolic blood pressure at 140 mm Hg; mean HbA1C at 7.6%; and mean low-density lipoprotein at 137 mg/dL. Patient health care satisfaction was high.

Conclusion: This practice-based intervention to foster social support for chronic care management among diabetics had no significant impact on the targeted outcomes. (J Am Board Fam Med 2008;21: 269–281.)

Major cardiovascular risk factors among Americans are highly prevalent but poorly controlled, especially among diabetics. Only one fourth of hypertensive Americans have their hypertension under control.^{1,2} The prevalence of dyslipidemia warranting treatment is approximately 29%, and control rates are probably no better than for hypertension.³ Approximately 20 million Americans have type 2

diabetes.⁴ Diabetes triples the risk of symptomatic cardiovascular disease (CVD), which causes two thirds of all deaths among diabetics.^{5,6} Morbidity and mortality are reduced when diabetics use lipid-lowering or antihypertensive therapy.^{2,5,6–8} However, the target levels for blood pressure (BP) and low-density lipoprotein (LDL) set for diabetics are especially difficult to achieve.^{2,7,8} Poor adherence to treatment and ineffective patient education often contribute to poor control of cardiovascular risk.^{9–12} Adherence to treatments is especially problematic for diabetics because of their complex treatment regimens.^{12–15}

Patients with diabetes typically obtain most of their medical care from primary care providers (PCPs).¹⁶ Practical strategies for use by PCPs to improve their patients' health behaviors are needed to realize the potential health impacts of powerful treatments. Patient education must be reinforced with other approaches for behavior change to have significant impacts on health.^{17–19} Social support is a key factor in the successful management of

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chronic diseases; it is a strong predictor for treatment adherence and favorable outcomes.^{9,15,20–22} However, most reported interventions used to enhance social support for chronic disease management are too costly and intensive to be feasible for most primary care practices under current reimbursement models. We therefore sought to evaluate the potential of a more practical primary care intervention to enhance social support for chronic disease management.

Potential pathways for social support to influence health outcomes are complex. Because health beliefs have been associated with health behaviors affecting cardiovascular risk,^{23–26} we used the Health Belief Model as a guiding conceptual framework and incorporated theories of self-efficacy.^{27–31} Self-efficacy (perceived health competence), the belief that one has influence over success and that one can succeed, has been positively associated with the likelihood of behavior change.²⁹ According to our model, cardiovascular risk factors such as hypertension and dyslipidemia are influenced by health behaviors such as adherence to prescribed diet and exercise, which in turn are influenced by health beliefs such as self-efficacy, perceived susceptibility, disease severity, and barriers. Personal relationships influence pertinent health beliefs. A support person (SP) can serve as a cue to action, reduce barriers to adherence, and promote self-efficacy; or, conversely, the SP might have detrimental effects on health behaviors through negative influences such as nagging.^{32,33} Moreover, the SP's influence may be moderated by factors such as the patient's age, gender, and education, the complexity of prescribed treatments, the patient's overall health status, and insurance coverage.^{9,34}

The purpose of this study was to test the effectiveness of an intervention to foster the involvement of a relative or friend as an SP in the control of cardiovascular risk factors in patients with type 2 diabetes. Our intervention was designed for broad and sustainable use in busy primary care practices. We avoided restricting the intervention to patients who had been formally assessed as most likely to benefit from such an intervention because we did not presume that such assessments would routinely occur in practice.

Methods

Details of our methodology have been published.³¹ This protocol was approved and monitored by the

Medical Institutional Review Board of the University of Kentucky.

Design Overview

We performed a clustered randomized controlled trial involving 18 primary care practices in the Kentucky Ambulatory Network, a practice-based research network (PBRN). The intervention was designed to educate, motivate, and facilitate patients and their SPs to work together to improve the patients' cardiovascular risk, health-related quality of life (HRQL), and satisfaction with health care. The roles of plausible mediating and moderating factors were evaluated, including medication adherence, basic relationship of patient to SP, quality and degree of SP involvement, patient demographics, history of CVD, health-related self-efficacy, and levels of social support. Participants had 9 to 12 months of follow-up. The study was restricted to patients willing and able to formally involve a SP in the control of the target conditions.

Settings/Locations

Each patient participant received care at one of 18 PBRN practices, where all study visits occurred. Participating PBRN practices were chosen based on their willingness to collaborate and their distance from the coordinating center. All data were collected at the primary care practices or via telephone. The study was coordinated at the University of Kentucky by Kentucky Ambulatory Network staff.

Patient Eligibility Criteria

1. Either type 2 diabetes based on chart review according to diagnostic criteria of the American Diabetes Association³⁵ or the diagnosis of type 2 diabetes recorded by the PCP along with a HbA1C level $\geq 8.0\%$, a random serum glucose level >200 mg/dL, or a current prescription for an antidiabetic drug.
2. Hypertension with suboptimal control, with or without uncontrolled dyslipidemia (see below).
3. Prepared to designate an SP with whom the patient would be in contact for the next 12 months.
4. At least 21 years old and able to give informed consent.
5. Not pregnant or planning to become pregnant within the next 12 months.

6. Planning to be available for follow-up for at least the next 12 months.

We defined hypertension with suboptimal control in diabetics as a mean systolic BP (SBP) >129 mm Hg based on the last 2 visits recorded in the chart, or on the highest 2 readings in the last 6 months, *and* a mean SBP >129 mm Hg based on 3 standardized readings taken at the screening study visit. (SBP was used as the sole BP criterion to facilitate hypothesis-testing and sample-size calculations.)

We defined dyslipidemia with suboptimal control in diabetics as LDL >100 mg/dL *and* confirmation of suboptimal control based on fasting lipid profile done at baseline. (LDL was used as the sole lipid criterion to facilitate hypothesis-testing and estimation of required sample size.)

These definitions for suboptimal control of hypertension and dyslipidemia match the recommended thresholds for starting or altering drug therapy in diabetics.^{2,5,36}

Support Person (SP) Eligibility Criteria

1. Adult able to give informed consent.
2. Regular (at least weekly) contact with the patient expected throughout the coming 12 months.
3. Chosen by the patient based on advice for effective SP provided by the investigators (can arrange at least weekly contact, SP willing to come to study visits, likely to be supportive of patient in improving health behaviors).

Identification and Enrollment of Participants

Potential participants were identified using each practice's billing data, followed by medical record review. To minimize selection bias, patients were not recruited during visits to the PCP. Each practice provided a list of all adult patients who had made at least one visit in the past 2 years and who had ever had diabetes coded as a diagnosis. Study personnel then reviewed all of these patient charts for eligibility. Potentially eligible patients were sent an explanatory letter about the study and signed by the PCP. During follow-up recruiting by telephone, study personnel worked down a randomly ordered list of these letter recipients until the a priori target number of 25 patients from each practice had been scheduled for a screening visit or until repeated attempts had been made to reach all patients on the list. Each patient participant was re-

quired to name a potential SP and all were instructed on criteria for choosing an SP. Study personnel enrolled the designated SP for each patient participant randomized to the intervention group. The SPs designated by patients randomized to the control group were not contacted by study personnel.

Randomization Methods and Study Groups

To avoid contamination, randomization was done at the practice level, and all participants at a given practice were assigned to the same treatment group. The project statistician maintained sealed envelopes containing group assignment. Masking and blocked randomization were used to prevent recruitment bias and unbalanced allocation. All patient participants at each practice were enrolled before group assignment was unmasked. Once group assignment was revealed, each patient was informed by phone; if assigned to involve a SP, the patient was instructed to bring the SP to the next study visit, which was the actual intervention visit.

Subjects were randomized to the control group or one of 2 intervention groups (A and B). Our design had 2 intervention groups to facilitate exploration of mechanisms and predictors of the intervention's impacts while minimizing the Hawthorne Effect. Participants in intervention group B had more contact with study personnel, who collected data to more deeply probe the mechanisms through which involvement of an SP might affect outcomes, but they did not receive any additional education or coaching. We planned to test our quantitative outcomes-related hypotheses through comparisons of the control group with intervention group A.

Sample Size

Our a priori sample size calculations were based on the plan to test our main hypotheses by comparing the control group with intervention group A. We used published variance data on SBP, LDL cholesterol, and HRQL scores to support sample size calculations.

Hypotheses Used to Drive Sample Size Calculations

After the intervention and controlling for baseline values, the following were hypotheses used to drive sample size calculations:

- Mean SBP will be at least 8 mm Hg lower in the intervention group than in the control group.
- Mean LDL cholesterol will be at least 15 mg/dL lower in the intervention group than in the control group.
- HRQL, as measured by the Medical Outcomes Study Short Form (SF-36) physical composite score, will be at least 10 points better (on a 100-point scale) in the intervention group than in the control group.
- We set our targeted sample size to detect at least these differences between groups with at least 90% power and an α level of 0.05. These calculations led to a targeted sample size of 100 patients per group completing the study.

Timelines

The entire study lasted 4 years (2002–2006). Participants at the first 15 practices were followed for 12 months; those from the last 3 practice sites had 9 months of follow-up.

Intervention

The intervention was designed to foster the formal involvement of a friend or relative as an SP to help the patient lower cardiovascular risk. It consisted of one patient/SP education session followed by 4 quarterly “newsletters.” Patients in the intervention groups had their SP join them for a 30-minute individualized patient education session with a Registered Nurse patient educator, delivered at the practice site. The session focused on cardiovascular risk reduction advice for the patient and ways that the SP could help. Guidelines for effective patient SP interactions were included, stressing the SP role as *facilitator* and advising that the patient should be responsible for his/her own health behaviors. Specific strategies suggested to the SP included accompanying the patient during doctor visits, reviewing medication instructions and supplies with the patient, exercising with the patient, grocery shopping with the patient, and talking with the patient about his or her concerns and specific barriers to cardiovascular risk factor control. After the education session, 4 quarterly patient education newsletters about cardiovascular risk factor control were mailed to the patients; similar newsletters focusing on facilitative strategies were mailed to the SPs.

Patients randomized to the control group received an individual 30-minute patient education

session with a Registered Nurse patient educator and received the same 4 patient newsletters as sent to intervention group patients, but control group patients did *not* have formal involvement of their SP in the program.

All education sessions followed a standardized curriculum guided by a notebook given to the participants. Notebook materials came from the American Heart Association, the National Institutes of Health, the American Diabetes Association, and the American Academy of Family Physicians, and were selected for accuracy, clarity, brevity, reading level at or below 8th grade, appropriate focus, and avoidance of confusing or conflicting recommendations. Information about CVD among diabetics, hypertension, high cholesterol, diet and exercise guidelines with examples, the importance of medication adherence and tips for improving it, advice on communicating with health care providers and keeping appointments, and advice on making lifestyle changes related to tobacco avoidance, diet, and exercise was included in the notebook. Intervention group materials included advice on how to help someone else make lasting improvements in their health behaviors. The newsletters reinforced the same curriculum, with each newsletter focusing on one or 2 of these topics.

Participants’ PCPs were aware of which patients were enrolled because they received study-related laboratory test reports (the same tests, regardless of study group assignment), but our patient-oriented intervention did not include any instructions or facilitation for the PCPs related to evaluation or management of cardiovascular risk.

Study Visit Protocol

All visits were performed at the practice sites. Each patient participant made 7 study visits over a 12-month period: 3 visits at baseline, 2 visits at 6 months after randomization, and 2 visits at 12 months after randomization. In intervention group A, SPs accompanied their patients to one visit at baseline and one at 12 months. SPs in intervention group B made one visit at baseline, one at 6 months, and one at 12 months. Patient and SP education activities occurred during visit 3. All other visits were strictly for data collection. Paired visits allowed repeated BP and LDL measurements to dampen intraperson short-term variability.

Table 1. Data Collected for Each Participant, by Study Visit

	Visit 1	Visit 2	Visit 3	Visit 4	Visit 5	Visit 6	Visit 7
Time since randomization* (months)	Baseline	Baseline	Baseline	6	6	12	12
Resting BP	A, B, C	A, B, C	A, B, C	A, B, C	A, B, C	A, B, C	A, B, C
Fasting lipid profile		A, B, C	A, B, C	A, B, C	A, B, C	A, B, C	A, B, C
HbA1C		A, B, C		A, B, C		A, B, C	
CVD history	A, B, C						
Interval CVD history					A, B, C		A, B, C
Medication review	A, B, C				A, B, C		A, B, C
Medication adherence			A, B, C		B		A, B, C
HRQL	A, B, C				A, B, C		A, B, C
Healthcare satisfaction	A, B, C						A, B, C
Health-related self-efficacy	A, B, C				B		B
Social network	A, B, C				B		A, B, C
SP qualities/involvement	A, B, C				B		A, B, C
Demographics	A, B, C				A, B, C		A, B, C

Patient education session occurred during visit 3.

*Randomization occurred after visit 2.

BP, blood pressure; CVD, cardiovascular disease; HRQL, health-related quality of life; SP, support person; A, intervention group A; B, intervention group B; C, control group.

Data Collection Protocol

The data collection protocol is shown in Table 1. Standardized BP readings and phlebotomy were performed by nurses and medical assistants employed at each practice. All blood samples were analyzed at one university laboratory. All other data were collected by study coordinators who went to each practice site. All nurses and medical assistants who took BP measurements completed training and certification in standardized BP measurement, following American Heart Association guidelines. Training and certification were repeated after 6 months. BP readings were done in triplicate to calculate average BP for that visit. LDL cholesterol

levels were collected as paired samples on different days to derive average levels for baseline and follow-up periods and were limited to the subset of patients whose most recent LDL on record was elevated. Participants with no LDL level on record did not have an LDL level drawn for this study. Instruments used to collect all other data are shown in Table 2.

Analytic Methods

Intervention group differences in baseline characteristics were assessed using mixed effects models where the clinic was treated as a random effect with the patient nested within the clinic, adjusting for

Table 2. Method and/or Instrument(s) Used for Non-physiologic Measures

Measure	Method(s)/Instrument(s)
CVD history	Patient structured interview with research staff
Interval CVD history	Patient structured interview with research staff
Medication review	Patient structured interview with research staff
Medication adherence	Medication Adherence Questionnaire ³⁷
HRQL	SF-36 Health Survey ^{38,39}
Healthcare satisfaction	Patient Healthcare Satisfaction Survey ⁴⁰ over telephone
Self-efficacy	Perceived Health Competence Scale ⁴¹
Social support network	Self-report at baseline on whether person(s) already helping patient with CVD risk management and person patient would ask to be SP (friend vs. relative)
SP qualities/involvement	Social Support for Intervention Survey ³²
Demographics	Patient structured interview with research staff

CVD, cardiovascular disease; HRQL, health-related quality of life; SP, support person.

the intraclass correlation coefficient (ICC) within the clinic.⁴² The ICCs for study outcomes calculated across the 18 clinic sites ranged from 0.0108 (satisfaction with health care) to 0.1046 (SBP).

Baseline characteristics and outcomes did not differ between the 2 intervention groups (A and B). The only differences between these 2 groups were in assessment, not intervention (see Table 1). Extra surveys were administered in group B to probe the mechanisms through which involvement of an SP might affect outcomes, but these participants did not receive any extra education, encouragement, or coaching compared with group A. To improve study power, we therefore collapsed groups A and B into a single intervention group for comparisons to the control group.

Main models to assess for differences between the intervention and control groups were fit to each individual outcome, adjusting for baseline values and ICCs. A second set of outcome models included adjustments for age, race, sex, employment, education level, health insurance, CVD event composite score, medication adherence level, perceived health competence, alcohol use, current smoking status, level of health-related social support, relationship with SP, and self-reported histories of high BP and high cholesterol.

The Statistical Analysis System procedure PROC MIXED (SAS Institute, Inc., Cary NC) was used to assess intervention effects on the outcome data while accounting for missing outcome measures because of drop-outs or sporadically missing data. Post hoc power calculations were conducted (see Results, below).

Results

Participant Flow and Baseline Characteristics

We enrolled 199 diabetic patients and 108 SPs from among 18 primary care practices and followed them for 9 to 12 months after randomization. Medical records on 2608 diabetic adults were screened; 1318 met initial eligibility criteria, 336 patients completed a screening visit, 233 patients were consented and enrolled, and 199 patients plus 108 consenting SPs completed the education session. The 34 patients “lost” before completing their education session included 4 who decided after consenting that they could not bring an SP and 30 who dropped out before completing the education session.

Baseline characteristics of the 199 patients are shown in Table 3. Randomization resulted in even distribution of values across the study groups; however, health insurance status differed: the control group had more privately insured patients (70%) than intervention group A (53%) or group B (52%), with group A having the most uninsured patients (14%). In addition, more patients reported having high cholesterol in the control group (80%) than in the intervention groups A or B (67%, 55%). SP characteristics are shown in Table 4.

Given the absence of significant baseline differences between intervention groups A and B, we pooled these into a single intervention group for our analyses; this allowed us to achieve sufficient power levels for all main outcomes except LDL cholesterol level.

Main Outcomes

Table 5 shows the main outcomes at 6 months and 9 to 12 months after randomization. For each outcome compared across groups, 2 *P* values are given. The first is adjusted only for baseline value and the ICC. The second *P* value is also adjusted for potential mediating or moderating factors (see Table 5 footnotes). Diminishing sample sizes over time reflect the drop-out rates.

Effects of Intervention

SBP fell in control and intervention groups, probably partly because of regression to the mean. The patient education received by all groups may have also played a role. There were no significant between-group differences in the change in SBP from baseline to 6 months and 9 to 12 months after randomization. There was also no significant effect of group assignment on diastolic BP (not shown).

LDL fell in the control and intervention groups, also probably partly because of regression to the mean. The patient education received by all groups may have also played a role. There were no statistically significant between-group differences in the change in LDL.

There were no significant between-group differences in the change in HbA1c from baseline to 9 to 12 months after randomization. There was a trend toward a slightly greater reduction in HbA1C levels in the control group at 6 months.

There were no significant between-group differences in the change in the Physical Composite Scale or the Mental Composite Scale of the SF-36

Table 3. Patient Participant Baseline Characteristics and Values

	Total Sample	Group A	Group B	Groups A and B Combined (AB)	Group C (Control)	P (A vs B vs C)	P (AB vs C)
N	199*	50	58	108	91		
Age in years (mean [SD])	62.1 (10.79)	60.3 (9.44)	62.0 (11.51)	61.2 (10.59)	63.1 (10.98)	.4400	.2644
Gender (%)							
Female	55.3	48.0	65.5	57.4	52.8	.1521	.5678
Race (%)							
White	86.9	88.0	82.8	85.2	89.0	.5257	.5279
African-American	13.1	12.0	17.2	14.8	11.0		
Employment (%)							
Employed	37.5	47.9	35.2	41.2	33.3	.3176	.2301
Retired	47.9	37.5	46.3	42.2	54.4		
Unemployed/disabled	14.6	14.6	18.5	16.7	12.3		
Education (%)							
≤ Some high school	16.6	20.0	13.8	16.7	16.5	.9030	.9663
High school grad/GED	41.2	44.0	39.7	41.7	40.7		
2-year degree/some college	22.6	16.0	25.9	21.3	24.2		
≥ 4-year college graduate	19.6	20.0	20.7	20.4	18.7		
Health insurance (%)							
Group/private	60.9	53.1	51.9	52.5	70.3	.0046	.0303
Medicaid/Medicare	32.8	32.7	42.3	37.6	27.5		
Other	1.0	0.0	3.7	2.0	0.0		
None	5.2	14.3	1.9	7.9	2.2		
Has current SP (% Yes)	29.7	26.0	34.5	30.6	28.6	.6006	.8763
Planned SP type (%)							
Relative	82.8	85.7	79.3	82.2	83.5	.6630	.8520
Friend	17.2	14.3	20.7	17.8	16.5		
Importance of SP help (%)							
Very	56.4	59.2	61.4	60.4	48.0	.1800	.2313
Somewhat	25.6	28.6	15.8	21.7	34.0		
Not really	18.0	12.2	22.8	17.9	18.0		
SP was helpful or bothersome							
Bothersome	7.0	12.2	3.5	7.6	5.9	.2212	.2796
No effect	10.2	6.1	8.8	7.6	15.7		
Helpful	82.8	81.6	87.7	84.8	78.4		

CVD event score (mean [SD]) [†]	0.95 (1.47)	0.74 (1.21)	0.93 (1.63)	0.84 (1.45)	1.08 (1.50)	.3433	.1724
History of high BP (%)							
Yes	86.9	84.0	89.7	87.0	86.7	.6843	1.0000
History of high cholesterol (%)							
Yes	69.9	55.1	66.7	61.3	80.0	.0076	.0050
Current smoker (%)							
Yes	15.6	16.0	8.6	12.0	19.8	.1861	.1698
Alcohol Use (%)							
Current use	28.4	22.0	32.8	27.8	29.2	.2821	.6264
Never/past use	71.6	78.0	67.2	72.2	70.8		
SBP (mean [SD])	140.3 (10.9)	142.7 (11.43)	140.2 (10.44)	141.3 (10.93)	139.0 (10.78)	.6771	.5433
Diastolic BP (mean [SD])	76.8 (9.00)	80.7 (8.26)	74.8 (9.20)	77.5 (9.23)	75.9 (8.68)	.1491	.4822
SF-36 physical subscale (mean [SD])	39.3 (11.16)	36.5 (11.30)	39.2 (11.65)	38.0 (11.51)	40.9 (10.55)	.1269	.0829
SF-36 mental subscale (mean [SD])	46.8 (11.94)	47.3 (12.75)	46.4 (12.03)	46.8 (12.31)	46.8 (11.54)	.9629	.9779
HbA1c (mean [SD])	7.6 (1.51)	7.5 (1.64)	7.5 (1.56)	7.5 (1.59)	7.6 (1.40)	.6860	.4102
Rating of personal doctor (mean [SD])	9.2 (1.07)	9.2 (1.05)	9.3 (0.97)	9.3 (1.00)	9.2 (1.14)	.8743	.6931
Rating of overall health care (mean [SD])	9.0 (1.33)	9.0 (1.55)	9.1 (1.11)	9.1 (1.32)	8.9 (1.37)	.6285	.3881
Perceived health competence (mean [SD])	3.2 (0.81)	3.3 (0.79)	3.3 (0.76)	3.3 (0.77)	3.2 (0.85)	.5539	.2278
Medication adherence (%)							
High	40.4	50.0	29.8	39.3	41.8	.1584	.4358
Medium	51.5	42.0	63.2	53.3	49.5		
Low	8.1	8.0	7.0	7.5	8.8		
LDL cholesterol							
N [‡]	40	8	16	24	16	.3037	.9471
Mean (SD)	137.1 (23.77)	127.1 (25.31)	142.0 (22.32)	137.0 (23.90)	137.3 (24.35)		

*N = 184–199; total sample = 199.

[†]CVD events score: 1 point each for past myocardial infarction, stroke, transient ischemic attack, coronary revascularization, and congestive heart failure.[‡]Number of participants in lipid measurement is only one fifth of total sample.

SP, support person; CVD, cardiovascular disease; BP, blood pressure; LDL, low-density lipoprotein.

Table 4. Support Person Characteristics

Age in Years (mean [SD])	55 (14.5)
Female (%)	75
Race (%)	
White	86
African-American	13
Asian	1
Occupation status (%)	
Employed	48
Retired	34
Unemployed	18
Education Level (%)	
<High school	17
High school grad or GED	35
Some college	25
≥4-year degree	23
Relationship to patient (%)	
Spouse	51
Child	16
Mother	5
Other relative	11
Friend and/or neighbor	17

from baseline to 6 months or 9 to 12 months after randomization.

Satisfaction with the PCP improved slightly in both groups, whereas satisfaction with health care overall declined slightly in both groups. Between-group differences became statistically nonsignificant after the described adjustments, and were not clinically significant on a scale of 1 to 10.

Power Considerations for Main Outcomes

Recruitment lagged behind our design target of 300 completed patient studies, leaving this study with marginal power to avoid type II errors in some of our outcomes analyses, and no power to detect differences in LDL levels. Ultimately, we had 80% power ($\alpha = 0.05$) to detect differences between the control group and the combined intervention groups (A and B) (see Table 6).

Potential Cognitive Behavioral Mechanisms

Linking the Intervention and the Main Outcomes

Two mechanisms were analyzed as potential moderators of the intervention effects on the main outcomes and as proximal outcomes themselves. Baseline levels of neither self-efficacy nor medication adherence were found to moderate intervention effects for any of the main outcomes. Furthermore, there were no significant between-group differ-

ences in the change in self-efficacy or medication adherence from baseline to 6 or 12 months after randomization.

Discussion

This randomized, controlled trial of a practice-based intervention intended to foster the formal involvement of a friend or relative in the care of adults with type 2 diabetes showed no significant effects on the main outcomes of interest, including SBP, LDL, HbA1C, HRQL, and patient satisfaction. The intervention also had no significant effect on perceived self-efficacy in managing one's own health or on medication adherence.

There are a few plausible explanations for this lack of impact. The first is that the intervention was not robust enough to bring about positive health behavior change beyond the effects of patient education. A single educational session with a nurse, followed by reinforcing newsletters, seems to be insufficient to foster social support at a level that leads to improved health behaviors. Second is the possibility of ceiling or floor effects within our sample. Although SBP >129 mm Hg and LDL >100 mg/dL determined eligibility for study entry, the mean levels of these risk factors in participants at baseline were not very high (140 mm Hg and 137 mg/dL, respectively). Having type 2 diabetes was an entry criterion, but HbA1C level was not, and the mean baseline HbA1C was only 7.6%. Thus, on the whole, these patients and their physicians may have felt less motivation for improving these cardiovascular risk factors than would be the case for patients with worse control. Furthermore, these patients' satisfaction with their PCPs and their overall health care was very high at baseline, leaving almost no room for improvement. Third, because this intervention required participants to name a potential SP, their overall social support may have been stronger than would be seen in the general population of adult diabetics; thus there may have been less room for improvement in support for cardiovascular risk reduction. We have no data to support or refute this possibility.

Limitations

Lagging enrollment led to a smaller sample size than was planned and required us to shorten the follow-up period to 9 months for the last 42 patients. This resulted in insufficient power to mea-

Table 5. Primary Outcomes: Adjusted* and Unadjusted† Group Differences at 6 Months and 9 to 12 Months After Baseline

	Baseline (mean [n])	6 Months (mean [n])	9 to 12 Months (mean [n])	Change from Baseline to 6 Months	Change from Baseline to 9 to 12 Months
Systolic BP (mmHg)					
Intervention group AB	141.3 (108)	135.5 (92)	134.0 (81)	−5.8	−7.3
Control group C	139.0 (91)	133.6 (74)	133.8 (60)	−6.4	−5.2
Unadjusted <i>P</i> for AB vs C	.5433	.3836	.9427		
Adjusted <i>P</i> for AB vs C	na	.4969	.6475		
HbA1C (%)					
Intervention group AB	7.5 (106)	8.3 (87)	7.4 (74)	0.8	−0.1
Control group C	7.6 (85)	7.8 (63)	7.4 (63)	0.2	−0.2
Unadjusted <i>P</i> for AB vs C	.4102	.0567	.6440		
Adjusted <i>P</i> for AB vs C	na	.0429	.9164		
SF-36 Physical composite score‡					
Intervention group AB	38.0 (107)	42.7 (84)	41.4 (74)	4.7	3.4
Control group C	40.9 (88)	42.6 (74)	41.6 (72)	1.7	0.7
Unadjusted <i>P</i> for AB vs C	.0829	.4145	.4345		
Adjusted <i>P</i> for AB vs C	na	.9598	.9056		
SF-36 mental composite score					
Intervention group AB	46.8 (107)	42.7 (84)	45.7 (74)	−4.1	−1.1
Control group C	46.8 (88)	40.1 (74)	47.9 (72)	−6.7	1.1
Unadjusted <i>P</i> for AB vs C	.9779	.2666	.5200		
Adjusted <i>P</i> for AB vs C	na	.2187	.2916		
Rate of primary doctor§					
Intervention group AB	9.3 (98)		9.5 (71)		0.2
Control group C	9.2 (86)		9.3 (67)		0.1
Unadjusted <i>P</i> for AB vs C	.6931		.0255		
Adjusted <i>P</i> for AB vs C	na		.6372		
Rating of overall health care§					
Intervention group AB	9.3 (98)		8.3 (71)		−1.0
Control group C	9.2 (86)		8.5 (67)		−0.7
Unadjusted <i>P</i> for AB vs C	.6931		.0255		
Adjusted <i>P</i> for AB vs C	na		.6709		
LDL cholesterol					
Intervention group AB	137.0 (24)	139.4 (18)	135.4 (18)	2.4	−1.6
Control group C	137.3 (16)	130.5 (11)	110.6 (11)	−6.6	−26.7
Unadjusted <i>P</i> for AB vs C	.9471	.6716	.3238		
Adjusted <i>P</i> for AB vs C					

*Unadjusted: *P* adjusted only for clustering of patient within clinic and baseline outcome variable values for 6 months and 12 month comparisons.

†Adjusted: *P* adjusted for clustering, baseline outcome values, age, sex, race, education, employment status, health insurance, whether already had an SP, relationship with SP, baseline values for medication adherence, perceived health competence, patient self-report of history of high BP, history of high cholesterol, smoking status, alcohol use and history of cardiovascular events score (1 point each for previous myocardial infarction, stroke, transient ischemic attack, coronary revascularization, and congestive heart failure). Given the limited sample size for LDL measurements, no adjusted *P* are presented.

‡SF-36 subscale possible range is 0–100.

§Doctor and healthcare satisfaction scales each have possible range from 0–10.

||The smaller subsample of patients with measured LDL permitted analyses controlling only for baseline LDL and clustering within clinic, not for adjusted analyses with additional covariates.

BP, blood pressure; LDL, low-density lipoprotein, SP, support person.

sure any significant effect on LDL and insufficient power to detect very small differences in other outcomes. Data were combined for participants

who were followed for 9 months and 12 months. Foreshortened follow-up of the last 42 subjects might be expected to falsely skew the results toward

Table 6. Minimum Detectable Between-Group Differences in Outcomes

Outcome	Detectable Difference at 6 mo	Detectable Difference at 9 to 12 mo
Systolic BP	10 mmHg	12 mmHg
LDL cholesterol	59 mg/dl	72 mg/dl
HbA1C (%)	1.3 percentage points	1.5 percentage points
SF-36 HRQL (100-pt scale)	6.5 points	6.9 points
Satisfaction with doctor (scale 0–10)	na	1.3 points
Satisfaction with healthcare (scale 0–10)	na	3.0 points

BP, blood pressure; LDL, low-density lipoprotein; HRQL, health-related quality of life.

a lasting intervention effect beyond 6 months, but we saw no effects at 9 to 12 months anyway. Non-random selection of participating practices might have led to unreliable results, but because this intervention was patient-oriented, not practice-oriented, it is unlikely that our results would differ if the study were repeated in a random sample of practices.

Conclusions

The prevention of symptomatic CVD is heavily dependent on health behaviors. Despite ample evidence that social support is positively associated with health and with health behaviors there is little published research about fostering social support to improve chronic disease management in the absence of an acute event, such as after a myocardial infarction. Our lack of impact with an approach that could be used in busy primary care practices is disappointing, but our study is importantly distinguished from those that used more intensive interventions that would be impractical for primary care practices. Other recent practical intervention trials^{43–45} have also failed to link fostering social support with positive health outcomes, such as improved glycemic control or serum cholesterol levels. As in our research, these studies focused on improving social support from family members and friends for the improvement of chronic disease management by using interventions that might be applicable in busy medical practices.

In contrast, studies showing favorable effects used more intense and/or prolonged social support interventions that focused on peer patient group visits,^{45,46} existing strong social support systems,⁴⁷ and other patient education and support group sessions.^{45,48} Interventions combining social support with diet, exercise, and stress management techniques had positive results.^{45,46,48} These interven-

tions included various combinations of weekly phone calls, weekly or monthly meetings, retreats, and regular group sessions with dietitians, exercise physiologists, nurses, and stress management specialists.

Practical and powerful methods that busy PCPs can use to foster sustained positive health behavior change in patients who have asymptomatic but dangerous chronic conditions are still needed. Even though the relatively successful approaches summarized above are not feasible for most primary care practices under current payment systems in the United States, PCPs may be able to guide the development of such programs and facilitate the involvement of their patients in them. Promising models that deserve further development and investigation are group visits for chronic care,^{49,50} improved integration of primary care services with work-place wellness programs and/or with chronic disease management programs offered by health insurers,⁵¹ and providing professional guidance to community programs, such as faith-based health improvement initiatives.⁵²

The most powerful solutions may lie in new models for health care reimbursement. Use of care teams coordinated through a primary care medical home and supported by a novel business model has shown promise for improving chronic disease management and deserves further investigation.⁵³ Any successful strategy for sustained improvements in such health behaviors may ultimately hinge on empowering, motivating, and equipping people to take greater proactive responsibility for their health.

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