Preparing Adolescents With Chronic Disease for Transition to Adult Care: A Technology Program

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KEY WORDS
transition to adulthood, adolescents, self-management, chronic disease management

ABBREVIATIONS
ACD—adolescents and young adults with chronic disease
CF—cystic fibrosis
IBD—intestinal bowel disease
SMS—short message service
T1D—type 1 diabetes
TRAQ—Transition Readiness Assessment Questionnaire

Dr Huang conceptualized and designed the study, obtained funding for the study, drafted the initial manuscript, and oversaw the entire project; Drs Gottschalk, Pian, and Norman reviewed the initial analyses and reviewed and revised the manuscript; Ms Terrones, Mr Tompane, and Ms Dillon designed the data collection instruments, coordinated data collection, and critically reviewed the manuscript; Dr Bartholomew worked with Dr Huang to conceptualize and develop the intervention and reviewed and revised the manuscript; and all authors approved the final manuscript as submitted.

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WHAT’S KNOWN ON THIS SUBJECT: Adolescents with chronic disease are a diverse population with common needs for transition. Disease-specific interventions have shown promise at improving patient outcomes but with substantial personnel and resource costs. Whether a generic approach across diseases may be useful is unknown.

WHAT THIS STUDY ADDS: This study is among the first to evaluate a generic (across disease) approach to transition of adolescents to adult care. The approach demonstrated promise and cost savings due to reduced personnel requirement and use of low-cost technology dissemination methods.

abstract

BACKGROUND: Adolescents with chronic disease (ACD) must develop independent disease self-management and learn to communicate effectively with their health care team to transition from pediatric to adult-oriented health care systems. Disease-specific interventions have been implemented to aid specific ACD groups through transition. A generic approach might be effective and cost-saving.

METHODS: Eighty-one ACD, aged 12 to 20 years, were recruited for a randomized clinical trial evaluating an 8-month transition intervention (MD2Me). MD2Me recipients received a 2-month intensive Web-based and text-delivered disease management and skill-based intervention followed by a 6-month review period. MD2Me recipients also had access to a texting algorithm for disease assessment and health care team contact. The intervention was applicable to adolescents with diverse chronic illnesses. Controls received mailed materials on general health topics. Disease management, health-related self-efficacy, and health assessments were performed at baseline and at 2 and 8 months. Frequency of patient-initiated communications was recorded over the study period. Outcomes were analyzed according to assigned treatment group over time.

RESULTS: MD2Me recipients demonstrated significant improvements in performance of disease management tasks, health-related self-efficacy, and patient-initiated communications compared with controls.

CONCLUSIONS: Outcomes in ACD improved significantly among recipients of a generic, technology-based intervention. Technology can deliver transition interventions to adolescents with diverse chronic illnesses, and a generic approach offers a cost-effective means of positively influencing transition outcomes. Further research is needed to determine whether improved short-term outcomes translate into an improved transition for ACD. Pediatrics 2014;133:e1639–e1646
Transition is the purposeful, planned transfer of adolescents with chronic medical conditions from child-centered to adult-oriented health care systems. The Institute of Medicine describes the health care transition for adolescents and young adults with chronic disease (ACD) as crucial to future health and psychosocial outcomes. Healthy People 2020 identifies transition planning as a major objective for ACD. However, there remain notable deficits in the transition experience for many ACD. Health care providers and researchers recognize that generic disease management skills are required for and may be effectively and efficiently addressed across all chronic diseases. Yet, to date, interventions that prepare adolescents for transition have often been disease-focused and delivered in-person, increasing cost and decreasing availability and efficiency. Such clinic-based interventions do not have optimal reach because they do not account for the poor engagement of ACD with the health care system.

In a randomized clinical trial, we evaluated whether a generic, Internet- and mobile phone–delivered disease management intervention would improve disease management, self-efficacy, and communication outcomes in ACD. Inflammatory bowel disease (IBD), cystic fibrosis (CF), and type 1 diabetes (T1D), with their diversity in clinical course and disease burden, were chosen to represent the broad disease spectrum in ACD. We hypothesized that a generic intervention designed to improve disease management, health-related self-efficacy, and patient-initiated communications would be more effective than a general health intervention in ACD for transition preparation.

METHODS

Eighty-one patients with IBD, CF, or T1D and aged 12 to 22 years old without cognitive impairment were recruited (October 2010 to March 2011) from a tertiary care pediatric academic medical center serving ~1 million youth. The sample size was 80 to account for a 10% attrition rate. The institutional review boards at the University of California, San Diego, and Rady Children’s Hospital approved the study protocol. Informed consent and assent were obtained before the study procedures.

**Intervention Groups**

Random assignment to the intervention group (MD2Me) or control groups (Fig 1; by random block assignment stratified by disease and generated by study statistician) occurred after the baseline visit. Participants were not blinded to study group assignment. Cell phones and plans were provided to MD2Me subjects who did not have a cell phone.

**MD2Me**

MD2Me subjects received an 8-month technology-based disease management intervention based on Bandura’s Social Cognitive Theory. The intervention targeted the self-management constructs of monitoring disease symptoms, responding to monitoring with appropriate treatments, and actively working with health care providers to manage care. For 2 months, subjects were asked to log in to a secure Web site weekly to receive theme-based materials outlining common disease management and communication skills, and lifestyle tips (See Supplementary Materials and Figures for more information). Disease-specific case studies were provided to increase usability. Tailored short message service (SMS) messages and queries were delivered (3–5 messages/week) to ensure that participants received and understood intervention messages. After 2 months, Web site access was provided as a disease management and information resource. Weekly reminder SMS messages were also delivered to reinforce previously introduced concepts and skills.

To facilitate patient-initiated communications, MD2Me recipients were given access to an automated SMS algorithm (Fig 2) that provided disease management decision support and a health care team communications portal. Subjects could activate the SMS to...
report health concerns. According to level of urgency (ie, nonurgent [to be addressed within 48 to 72 hours] or urgent [requiring same-day attention]), subject concerns were relayed to the health care team (by page or e-mail for intervention). Usual health care communication portals (eg, phone) were available to all participants.

**Controls**

Controls received monthly messages via mail or e-mail (participant preference) addressing general health issues. Disease-specific information was provided as appropriate (eg, in the healthy nutrition module). Usual health care communication portals (eg, phone) were available to all participants.

**Assessments**

Participants’ disease self-management and health-related self-efficacy were assessed at study visits at a clinical research office by measurement staff blinded to group assignment. Patient-initiated communications were assessed as noted below. Health literacy, health status, and demographic data were also collected. All instruments have been used in adolescents and found to be reliable and valid.

**Disease Status**

Disease status was assessed by using scales developed for each disease. We used the Pediatric Ulcerative Colitis Activity Index (range: 0–85, with 0 indicating no disease and higher scores indicating worse disease)7 for patients with ulcerative/indeterminate colitis and the abbreviated Pediatric Crohn’s Disease Activity Index (range: 0–70, with 0 indicating no disease)8 for patients with Crohn disease. For CF, we used the Cystic Fibrosis Clinical Score (range: 10–50, with 10 indicating no disease)9. The Diabetes Quality of Life Brief Clinical Inventory (range: 60–0, with 60 indicating perfect disease status satisfaction)10 was used for T1D. Baseline predicted forced expiratory volume in 1 second and glycohemoglobin were extracted from the medical record for patients with CF and T1D, respectively.

**Global Health Status**

The Karnofsky Performance Scale,11 a functional status assessment scale used in chronic disease populations (0–100, where 100 represents perfect health and 0 represents death), was used as a universal functional status measure across all diseases. We also used the Pediatric Quality of Life Scale (PedsQL) to measure quality of life as an indicator of health status.12

**Health Literacy Assessments**

Health literacy was assessed by using the Test of Functional Health Literacy in Adults.13 The adult version was used to reflect the health information processing needs of patients transitioning to adult health care that was still sensitive to adolescent concerns.14,15 The Test of Functional Health Literacy in Adults provides information on both reading comprehension and numerical ability, with higher scores (0–100 range) indicating better literacy. Scores of <60 represent inadequate literacy.13

**Disease Management and Self-efficacy**

The Transition Readiness Assessment Questionnaire (TRAQ) 4.116 is a measure of readiness for transition and assesses performance of chronic disease self-management skills. The TRAQ 4.1 was scored by using a 6-point scale where 0 points were given for subjects who felt that the skill was not necessary for their care, 1 point was given for not knowing how to perform the necessary skill, and 5 was given for performing the skill consistently when needed. The TRAQ assumes that subjects can correctly identify when a skill is not necessary for their care. To better capture the inability of subjects to recognize that a particular skill was needed for disease self-management, we scored inability to recognize a skill as needed for chronic disease self-management as −1; the scale was otherwise scored the same. In cases in

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**FIGURE 2**

MD2Me SMS algorithm decision tree. Nonurgent, requiring intervention within 48 to 72 hours; Urgent, requiring intervention within 24 hours.
which a given skill was deemed not necessary for chronic disease self-management for a given patient situation, 0 points was assigned to the question. Because none of our patients was financially independent, all of the financial capability questions in TRAQ 4.1 were scored 0 points. A TRAQ summary score $\geq 4$ is equivalent to respondents reporting that they are starting to perform necessary disease management skills.

The Patient Activation Measure\textsuperscript{17} gauges participants’ self-efficacy and confidence essential to managing one’s own health and health care on a scale of 0 to 100, with 100 representing ideal health. Intervention outcomes (disease management, health-related self-efficacy, and patient-initiated communications) were analyzed by intent-to-treat randomization assignment, using repeated-measures models testing the treatment $\times$ time interaction. Variables expected to influence outcomes such as age, specific disease (ie, CF, T1D, and IBD), and literacy status were entered into outcome analyses. Statistical analyses were performed by using JMP statistical software (SAS Institute, Cary, NC); significance was set at $P < .05$.

RESULTS

Eighty-one youth with chronic disease were randomly assigned into the study (Fig 1). Demographic data are presented in Table 1. The ages of subjects in each disease group differed (median [interquartile range] for IBD, T1D, and CF: 17 [16–18], 17 [16–18], and 14 [13–16] years, respectively; $P = .004$). There were no significant ethnic or gender differences. Seventy-five youth completed the study. There were no differences between the 6 dropouts (2 MD2Me subjects and 4 controls) and the main cohort by age ($P = .20$), disease ($P = .10$), gender ($P = .53$), or ethnicity ($P = .10$). There were no

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<tr>
<th>TABLE 1</th>
<th>Demographic and Other Information on Study Participants</th>
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<tr>
<td>Characteristic</td>
<td>MD2Me ($N = 40$)</td>
</tr>
<tr>
<td>Age, median (minimum, maximum), y</td>
<td>17 (12, 20)</td>
</tr>
<tr>
<td>Gender (male-female), n</td>
<td>17:23</td>
</tr>
<tr>
<td>Race/ethnicity, n</td>
<td></td>
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<tr>
<td>White</td>
<td>16</td>
</tr>
<tr>
<td>Black</td>
<td>4</td>
</tr>
<tr>
<td>Hispanic</td>
<td>18</td>
</tr>
<tr>
<td>American Indian/Alaskan</td>
<td>1</td>
</tr>
<tr>
<td>Other</td>
<td>1</td>
</tr>
<tr>
<td>Disease category, n</td>
<td></td>
</tr>
<tr>
<td>IBD</td>
<td></td>
</tr>
<tr>
<td>CD</td>
<td>11</td>
</tr>
<tr>
<td>UC</td>
<td>6</td>
</tr>
<tr>
<td>CF</td>
<td>6</td>
</tr>
<tr>
<td>T1D</td>
<td>17</td>
</tr>
<tr>
<td>Disease duration, median (IQR), y</td>
<td>4 (3–9)</td>
</tr>
<tr>
<td>Disease status, median (IQR)</td>
<td></td>
</tr>
<tr>
<td>UC</td>
<td></td>
</tr>
<tr>
<td>PUCAI</td>
<td>3 (0–21)</td>
</tr>
<tr>
<td>CD</td>
<td></td>
</tr>
<tr>
<td>PCDAI</td>
<td>5 (5–10)</td>
</tr>
<tr>
<td>CF</td>
<td></td>
</tr>
<tr>
<td>CFCS</td>
<td>19 (17–24)</td>
</tr>
<tr>
<td>FEV\textsubscript{1%}</td>
<td>87 (69–94)</td>
</tr>
<tr>
<td>T1D</td>
<td></td>
</tr>
<tr>
<td>DQoL</td>
<td>47 (44–49)</td>
</tr>
<tr>
<td>Glycemic hemoglobin, %</td>
<td>8.2 (7.9–9.2)</td>
</tr>
<tr>
<td>Karnofsky Performance Scale, median (IQR)</td>
<td>90 (80–100)</td>
</tr>
<tr>
<td>TOFILA, median (minimum, maximum)</td>
<td>82 (74, 88)</td>
</tr>
<tr>
<td>PedsQL, median (IQR)</td>
<td>81 (70–91)</td>
</tr>
</tbody>
</table>

CD, Crohn disease; CFCS, Cystic Fibrosis Clinical Score; DQoL, Diabetes Quality of Life Brief Clinical Inventory; FEV\textsubscript{1%}, baseline predicted forced expiratory volume in 1 second by percentage; IBD, inflammatory bowel disease; IQR, interquartile range; PCDAI, Pediatric Crohn’s Disease Activity Index; PedsQL, Pediatric Quality of Life Scale; PUCAI, Pediatric Ulcerative Colitis Activity Index; TOFILA, Test of Functional Health Literacy in Adults; UC, ulcerative colitis.
study-related adverse events or confidentiality breaches.

**Disease Status**

Disease status as measured by specific disease scales ($P = .18–.86$), functional performance ($P = .31$), and quality of life ($P = .17$) did not differ according to treatment group assignment (Table 1). Disease severity varied by disease, as expected (Table 1).

**Baseline Health Literacy and Readiness for Transition (Disease Management and Health Self-efficacy) Measures**

Significant baseline health literacy, disease management, and health-related self-efficacy differences were noted across disease groups (Table 2) but not according to treatment group (Table 3). Most patients demonstrated adequate health literacy (74%). In contrast, only 34% (26% of MD2Me subjects and 41% of controls, $P = .13$) had begun to perform the necessary disease management skills needed for transition (defined as TRAQ $\geq 4$). Both treatment groups had minimal patient-initiated outside-of-clinic-visit contact with their health care teams at baseline.

**Change in Chronic Disease Management and Patient-Initiated Communications Outcomes Over Time According to Treatment Group**

MD2Me recipients demonstrated significant improvements in all targeted outcomes across the study period compared with controls, controlling for age, baseline literacy status, and disease group (Table 3). Specifically, MD2Me participants, compared with controls, demonstrated increases in disease self-management and health-related self-efficacy. In addition, the number of patient-initiated communications between MD2Me participants and their health care team increased significantly compared with the control group. This increase in patient-initiated communications by MD2Me participants primarily occurred via the SMS algorithm (median [minimum, maximum] increased interactions during study period by MD2Me participants: +1 (0, 7) SMS vs. +0 (0, 2) phone).

**DISCUSSION**

We demonstrated the efficacy of a technology-based, transition intervention to improve disease management, health-related self-efficacy, and patient-initiated communications in ACD across 3 disease groups. Our findings suggest that a more resource-intensive disease-specific approach may not be necessary now that a generic approach to aid ACD through transition has been shown to be efficacious. The use of technology in our program offers a low-cost alternative for intervention in this at-risk population, which can be distributed across time and geography and independent of the clinic setting.

**MD2Me: Compliance and Performance**

Among MD2Me participants, 79% (63%, 93%) (median [interquartile range]) of curriculum receipt was documented across both Web and SMS modalities. Web visits occurred primarily during the initial 2 months (All but 1 participant visited the Web site during the initial 2 months, whereas only 16 [42%] users visited in the latter 6 months). In contrast, all but 1 user responded to SMS curricula throughout the intervention with 78% (56%, 90%) compliance. Seventy-seven percent of participants used the SMS algorithm, and SMS algorithm users were older than those who did not access the SMS algorithm (mean [SD]: 15 [2] vs 17 [2] years; SMS algorithm non-users v. users, $P = .01$).
The need for ACD to be supported and prepared for transition is well recognized. Consensus statements (both disease-specific and generic) on the needs of the adolescent during this period have been published. To prepare for the transition from child-centered to adult-centered care, ACD should learn the skills needed to effectively care for themselves and how to advocate for their own health issues. Consensus guidelines suggest that this process should begin by age 14 years among ACD.

A generic approach to readiness for transition programs has not often been used in pediatrics despite evidence that low-cost disease management training using a generic approach improves health outcomes among adults with chronic disease. We demonstrated successful use of a generic approach for transition preparation interventions in ACD. In fact, disease type did not appear to affect the intervention effect on any of the targeted outcomes in our study despite notable differences between disease groups at baseline in disease management, health literacy, and health-related self-efficacy. The approach to transition support tested in this study offers availability and utility across diseases. The technology-based approach offers the possibility of efficient implementation in a clinic setting with modest personnel time and effort.

Recent trends indicate a major shift to incorporate technologies into health behavior interventions. These technologies offer several advantages for health behavioral interventions, such as information and messages tailored to the participant, quick access, increased cultural sensitivity, and anonymity, which may be attractive regarding sensitive health issues. Similarly, mobile technologies can be used to improve health outcomes. The computing power and portability of mobile phones make possible new applications for automatic, timely, and tailored presentation of health messages, and because mobile phones are generally only accessed by a single individual, outreach can be improved for sensitive medical issues. Wireless devices are now ubiquitous, and text messaging is currently the main form of communication by adolescents today. We previously demonstrated notable interest among ACD in using technologies in a disease management intervention, which has since been validated by others. Studies using the Web and SMS for promoting disease self-management among youth have found some success. Similarly, we demonstrated efficacy of our Web- and SMS-based intervention for chronic disease management in ACD. Most MD2Me recipients reported that the program was helpful as delivered via text (89%) and Web (91%), with 92% reporting that they would likely recommend the program to their friends or family. Most notably, the SMS communication portal appeared to encourage increased communications between participants and their health care team. Although the absolute increase in number of communications between ACD and their health care team is relatively small, it signified an increase from no communication to some communication. Lack of health care engagement remains a major issue with ACD and the demonstration of ACDs beginning to initiate communications with their health care team is an important step toward autonomy in managing one’s own health. In 1 situation, this communication led to timely intervention in 1 participant who required urgent admission for intensive medical care. Although it is possible that the participant would have contacted his health care team through usual channels had the SMS portal not been available and that this patient’s course was not altered via this SMS-enabled communication, review of past patient-initiated communications revealed no previous participant-initiated communications despite severe disease symptoms.

Suboptimal health literacy has been linked to poorer chronic disease and health outcomes among adults. Among children, low parental health literacy also appears to adversely affect children’s health outcomes. Similarly, we found that adolescent/young
adult health literacy status affected the
effects of a transition intervention on
disease management outcomes in ACD.
Apparent, participants with lower
health literacy did not accrue all of the
benefits of the intervention. This find-
ing may indicate the need for additional
modules to bolster the adolescents’
capacity to make use of the program if
they screen below a certain threshold in
health literacy.
We did not find significant effects of age
on disease management, health-related
self-efficacy, or patient-initiated com-
munication outcomes. This finding is
potentially important because young
adolescents are not often given the
autonomy to make independent health
decisions due to their presumed im-
maturity. Although many guidelines
recommend that transition and disease
management training should begin by
age 14 years, others promote engaging
the child affected by chronic disease at
an even younger age\(^{35}\) to set up a cul-
ture of learning and engagement
throughout what will be a lifetime of
health care experiences. Our findings
support that benefit can be derived
from intervening during early adoles-
cence because self-efficacy and patient-
initiated communication improvements
were seen independent of age and literacy status in MD2Me-assigned
participants.

Limitations of our study include the
relatively small sample size, single-
institution recruitment, and specific
patient populations studied. Neverthe-
less, we recruited at the main pediatric
referral center for a catchment area of
\(~1\) million persons (<18 years) across
a broad variety of ACD. Although we
demonstrated success in our generic
intervention approach to prepare ACD
for the transition process, we only
tested it in 3 patient populations.
Nevertheless, these patient cohorts to-
gether represent a broad range in
health status, disease, and disease
management burden. Although we
would like to attribute improvements
in the treatment group to the MD2Me
intervention, an alternative explana-
tion for the improvements may have been the increased frequency of
intervention interactions in the MD2Me
weekly) compared with the control
(monthly) group. The relatively short
duration of the intervention and follow-
up period likely limited our ability to
affect disease and functional status.

Conclusions
We demonstrated efficacy of a generic
intervention approach to improve dis-
ease management skills, health-related
self-efficacy, and patient-initiated com-
munications in ACD. The use of a common
approach and available technologies
offers a potentially low-cost option
for a population that is sparse and
widely distributed. Further study is re-
quired to determine whether demon-
strated improvements translate into
an improved transition experience
for ACD.

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