Social Networking Services-Based Communicative Care for Patients with Diabetes Mellitus in Korea

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Keywords

Diabetes mellitus, personal health records, communication, networks, home health, patient self-care, home care and e-health, consumer health

Summary

Objectives: Social networking services (SNS)-based online communities are good examples of improving quality of care by incorporating information technology into medicine. Therefore, we created an SNS-based community care webpage for communication among patients with diabetes mellitus (DM). We aimed to identify what diabetic patients wanted to know and were interested in by analyzing their posts and classified content in which users were interested.

Methods: As opposed to the existing physician-focused health information websites, we built a patient-focused experience exchange website, "I love insulin (<http://www.iloveinsulin.co.kr>)." The DM communication webpage was divided into "My Web-Chart," "My community-free board," and "Life & Health." The contents analysis targeted users' postings, and replies were classified by theme from May 2012 to June 2013. The data included number of questions asked, answers, and question-to-answer (QA) ratio in each category.

Results: A total of 264 patients registered on the "I Love Insulin" website. The most frequent topic of posts classified as questions were about diabetes itself (23%), diet (22%), and glucose levels (19%). Conversely, most answers and information provided by users were about daily life with no relationship to diabetes mellitus (54%). While there were many questions about diet, there were very few answers (2%). Whereas there was much provision of knowledge about general DM, sharing diet information was rare. The ratios of answers to questions on diet (ratio=0.059, 1/17), glucose level (ratio=0.067, 1/15), insulin regulation (ratio=0.222, 2/9) and webpage (ratio=0.167, 1/6) were significantly low compared to DM itself (all $p < 0.001$).

Discussion: Patients in Korea with DM tend to have insufficient knowledge about diet and insulin regulation; continuously providing diet and insulin regulation information are desirable. It is hoped that the patients would be motivated to participate actively by "knowledge sharing." Through this process, patients learn about their diseases not from the physicians but from among themselves.

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1. Introduction

Internet users obtain health information more commonly by searching through web portals than from their physicians [1]. In the United States, about 70–80% of American Internet users search for health information [2]. In addition, health information obtained through the Internet helps the users in the actual decision-making process [3]. Recently, social networking services (SNS) are drawing attention as mediums of prompt health information. [4–9]. Because SNS enable direct communication among users without much knowledge about computers [10] and allow them to easily access health information, users can share their opinions bi-directionally without limitations of time and space and obtain a substantial amount of health information. Therefore, it would be beneficial if health information and health content were efficiently supplied via SNS, and active participation and lifestyle modification of Internet users (medical consumers) were encouraged.

Many websites and applications targeting patients with diabetes mellitus (DM) have been released recently, as this patient population is increasing rapidly [11–13]. In fact, potential blood glucose management and clinically positive effects of using information technology (IT), such as medical websites or apps are constantly being reported [14–17]. Many recent studies address communication between doctors and patients with the help of IT, [14–15, 18–21] and physicians unilaterally convey health information to the patients. Although medical consumers are gradually obtaining more knowledge, the incidence of chronic diseases is still increasing significantly [22]. Excess knowledge about exercise and diet is better than not having any knowledge at all. However, chronic diseases increase not because the patients do not know much about diet or exercise, but because they do not implement what they know [23–25]. Therefore, what really matters is actively helping people bring about behavioral changes rather than just teaching them about these changes [25–27]. This would help in motivating patients to adopt the right habits and regime for their disease management. This study attempts to bring about active behavioral changes in patients by providing them a platform for sharing their challenges and knowledge about blood glucose management. In view of this, it would not be effective for medical consumers to simply gain more knowledge – medical consumers need to be active in online communities when considering their management of health risk factors. There seems to be a need of a platform to share information among patients, not just between patients and doctors. In fact, there are many online communities about diabetes formed by the patients themselves, and not by doctors. However, patients in these communities mainly focus on sharing their personal stories rather than how to manage blood glucose levels. There is also a lot of pressure about accuracy and credibility [28] of the health information provided on these websites, besides its proper functioning. Moreover, there could be contents for advertisement or untested health information on these websites as well. In addition, analyzing information shared among patients about blood glucose management could allow us to identify the needs of patients in managing their blood glucose levels.

Therefore, the primary aim of this study is to develop an SNS-based community care webpage based on communication among patients with DM. We also classified and confirmed the content of posts on this site in which users (patients with DM) are interested or have questions based on their difficulties. By providing proper feedback for difficult situations or answers to questions from users, we hoped to be of help to patients. We also hope to change medical consumers into active participants in addition to providing an information system for basic health information and knowledge.

2. Materials and Methods

2.1 Establishment of DM Communication Webpage (<http://www.iloveinulin.co.kr>)

In contrast to the existing health information websites, we emphasized creating a patient-focused experience exchange website rather than a physician-focused one. “*I love insulin*,” the DM communication webpage we built, was divided into “*My Web-Chart*,” “*My community-free board*,” and “*Life & Health*” (► Figure 1). In “*My Web-Chart*,” users can record blood glucose management data. In “*Life & Health*,” physicians provide DM information to users. We added patient-patient bulletin

board “*My community-free board*” for the users to share their opinions and feelings for the purpose of this study. “*My community-free board*” enables users to communicate and share their experiences.

Patients with DM who receive insulin injections can record their own blood glucose levels and HbA1c values under the “*My Web-Chart*” menu. In this menu, various data are recorded with other patients’ average values for comparison. It enables the patients to compare with others using various searches and methods and to accurately understand their current blood glucose status. This is intended to motivate the members of the website. The users can find other patients with the same type of diabetes, blood glucose status, and health condition under the “*My community-free board*” menu where they can discuss the difficulty of blood glucose management and exchange useful information with one another. In the “*My community-free board*” menu, only users can post notes on the website; the medical staff members do not intervene in the conversations among patients. Users are able to check the comments and respond and can be registered as “*favorite friends*.” If users click on others’ profile pictures, they can find information on age, DM duration, hospital, and HbA1c values of other users. It also allows users to register another user as their favorite (► Figure 2). Online users could use keywords such as DM type, duration, or treatment type to search for others’ comments of interest. “*Life & Health*” comprises information about patients with DM and insulin treatment. In the “*Life and Health*” menu, physicians who have their own private hospitals can write a column or update medical opinions under the manager’s permission. To convey accurate and reliable information, patients cannot write but can only read the comments. However, it is possible to write a column about medical controversies and issues on the “*My community-free board*.” The medical staff checked posts on “*My community-free board*” each week, found out what users were interested in, and posted relevant posts on “*Life and Health*” Menu.

2.2 Privacy Protection

The promotion for the website was made for random visitors to the endocrine internal medicine department in our hospital, but the registration was made at home. The website is not for only appointed hospitals or any patient who is interested could participate. Advertising and marketing for the study was conducted by our hospital. To apply for membership, patients had to sign a consent form and conduct email verification; they did not have to provide their *national identification number*, patient hospital ID, or real name. The users can search for information on diabetes and related current information. Our study was approved by the institutional review board of the Catholic University of Korea.

2.3 Analysis of Online Users’ Postings

The content analysis targeted users’ postings on “*My community-free board*” and “*Life & Health*” from May 2012 to June 2013. Each post became an analysis unit and its contents were designated as a subject of analysis. The contents were classified by two aspects: purpose (question or answer) and theme (health care, diabetes, etc.). Posts about DM were classified by diet, exercise, and medical themes. Posts with more than two themes were classified by allowing for overlap. In other words, when two topics occurred in one post, two themes could exist. Analysis by word, character, space, and time did not occur; posts were classified by focusing on what users were trying to say. We made various topic categories about DM, education, etc. to classify the writing in advance before classification. To classify the topics, one physician performed the classification and the other physician reviewed it once more to satisfy objectivity and generality. After classification by theme, the number of replies in each theme was confirmed.

2.4 Statistical Analysis

The posts were separated by questions and answers and were re-categorized by topic. Checking the numbers by category, we calculated the number of answers per question (QA ratio). Categorical variables were analyzed by chi-square test or exact test. Chi square tests were performed by comparing DM QA ratio as a reference to other categories. All statistical analysis were done with SAS version 9.4 (SAS Inc., Cary, NC) and p-values less than 0.05 were considered statistically significant.

3. Results

3.1 Analysis of “My Web-Chart” Menu

A total of 264 patients registered for the “*I Love Insulin*” website. Out of these 264 patients, 140 provided personal information so the baseline characteristics were obtained based on that information (► Table 1). The patients comprised 97 male (69.3%) and 43 female (30.7%) users. The mean age was 40.7 ± 11.7 years; three were above 65 years old. The mean BMI was 27.4 ± 11.8 kg/m². Specifically, there were 23 patients with Type 1 DM, 85 patients with Type 2 DM, and 12 patients did not answer the question. Twenty patients (20/140, 14.3%) did not know their DM type. DM duration was 8.5 ± 7.4 years. Of them, 68 patients (68/140, 48.8%) used only insulin in managing blood glucose and 29 used insulin in combination with oral medication (29/140, 20.7%) for blood glucose control. The rest of the patients were on oral medication only. Some of them had an insulin injection before or were recommended by doctors to receive insulin treatment in the future. The average HbA1c was $8.1 \pm 1.7\%$.

3.2 Analysis of “My community-free board” Menu

Only 33 patients posted at least once. The total number of posts was 222 cases on 457 topics with an average of 42 views per post. When posts were classified by topic, talking about one's health state including blood glucose condition were the most frequent (18%, 82/457). Following in order were 77 questions (17%, 77/457) related to DM, 57 encouragements from other people (12%, 57/457), 57 answers for questions or information delivery (12%, 57/457), 56 weather-related articles (12%, 56/457), and 36 daily life chats (8%, 36/457). In addition, there were 30 cases of determination of blood glucose management (7%, 30/457) and 24 cases of self-reflection on recent blood glucose management (5%, 24/457; Figure 3). Average replies per post were 3.0 ± 2.1 . The number of replies for questions about insulin and diet had the highest averages at 4.7 ± 2.3 and 4.7 ± 2.2 respectively. The replies about exercise cures were 3.5 ± 1.4 . However, those replies were not exactly questions, but more sympathetic content.

The most frequent topic of posts classified as questions were about DM itself, diet, glucose levels, insulin regulation, and exercise (► Table 2). Conversely, most answers and information provided by users was about daily life with no relationship to DM and the contents on DM itself were about only a third of posts. Replies to other questions were below 7%. For example, even though questions about diet and glucose level were roughly a fifth of the content, answers to those questions or information provision were very low. The ratios of answers to questions on diet, glucose level, insulin regulation, and webpage were significantly lower than DM itself (all $p < 0.001$).

3.3 Analysis of “Life and Health” Menu

A total of 24 posts were written by doctors, nurses and nutritionists employed at Seoul St. Mary's Hospital in Korea. Post topics in the early stages were mainly instructions and insulin dosage regulation, fitting the purpose of this site. However, as posts included in “*My community-free board*” were checked monthly, contents that became issues between users or could not be answered were mainly covered. The average number of people who viewed the post of the total of 24 posts was 118 ± 39 and posts about how to regulate basal (265 views) and bolus insulin (176 views) dosages had the most views, consistent with the purpose of the site.

4. Discussion

This study aimed to develop a SNS-based community care webpage based on communication among patients with DM. We found that users talked about their blood glucose condition very often; posting many questions about it was the most frequent communication method. The mean BMI (27.4 ± 11.8 kg/m²) in our study was similar to that of a previous Korean study (25.1 ± 3.5 kg/m²) [29]. Thus, we assumed that many people who used this website were diabetic and obese.

Users were concerned about their health (especially diet, exercise, and weight control, etc.), possibly because they had both DM and obesity [30]. Communication on various topics progressed without a medical team in the role of providing an answer or related information. Self-reflection or determination/plan was referenced frequently so that it would contribute somewhat to patients or others changing their minds.

In posts, content about daily life not related to DM was frequent. Users with DM maintained communication with each other by talking about general topics, besides DM (Figure 3). We first thought that people would mostly have a certain degree of knowledge and information about diet and exercises. However, we found that while there were many questions about diet, there were very few answers. Moreover, though there was much provision of knowledge about general DM, sharing of diet information was infrequent. Therefore, we concluded that patients with DM had insufficient knowledge about diet and there was a need to continuously provide diet information. The low responses to blood glucose conditions or insulin control could have been due to difficulties in discussing blood glucose conditions among DM patients. Indeed, previous studies have found that diabetic patients do not really know much about their blood glucose or the medicines that they are taking [31-32]. Thus, even if they are getting insulin injections, they are not likely to know why they are taking them or how to adjust the amount (► Table 2). To rectify this lack of knowledge, the medical team constantly checked posts on the menu “*My community-free board*” and posted information on the menu “*Life & Health*” about user-generated issues or content on the menu “*My community-free board*.” With this, users expected to obtain better knowledge about DM using this site.

In order to encourage active behavioral change in patients with DM, various methods are suggested [33-34]. Currently, as SNS are utilized in health and medical services, patients are increasingly sharing their personal health-related information online [1]. Our site allows users to manage identities, have conversations, form relationships online, share information, and build reputations for people or products. This has not been achieved under the existing doctor-centered treatment system. This “*knowledge-sharing type*” of communication among patients is different from the existing “*knowledge-conveying type*” of communication between physicians and patients. The main service of our patient-to-patient communication presents a method to constantly “self-care” a healthy lifestyle by obtaining information on disease prevention and medical care. We wish this website could be a chance for them to share information and experience about their health condition and blood glucose management besides what is provided by a hospital. A big difference between our website and other online web communities is that in this study medical staffs continuously monitor the posts. This is possible because it's an online webpage operated by a medical team. As previously mentioned in the study, the medical staff continuously monitored postings on “*My community-free board*” menu on the website and deleted uncertain, untested information and advertisements. Thus, this website can be trusted and relied on by the patients. It is also helpful for the medical staff as they can find out the main interest of the patients, which is the biggest advantage of the website.

The website was made by medical staffs of a certain hospital, but it mainly focuses on sharing information among patients without direct contact between the medical staff and the patients. The only thing the medical staff does is posting common reliable information to answer the patients' questions in “*Life and Health*” menu, not “*My community-free board*” menu. There is minimum intervention of the medical staff limited to deleting uncertain or untested information. The study focused more on the patient-patient based online system unlike doctor-patient online based blood glucose management.

Even though it was a site for patients, our webpage was advantageous for doctors as well. As current doctors are only accustomed to treatment in a hospital, they lack practical experience and knowledge for outpatient care. The relationships between physicians and patients still strictly focus on treatment. Physicians now share much more information with patients and interpret it for them rather than perform diagnosis and treatment based on their medical knowledge and clinical experience when dealing with chronic disease. It will also be possible for patients to manage their health between consecutive visits to the hospital [35-37]. Personal and family history in individual health status need to be considered and individualized health information content (diet, exercise, etc.) need to be provided to connect hospital, home, and lifestyles. This form of communication has the benefit of requiring patients' active participation. If medical consumers are provided with objective and adequate information about their health status from the point of questioning their health status to the

point of visiting the hospital, they may be more effective in maintaining healthy lifestyles [38]. Thus, our website could go beyond forming bonds through sympathy and providing information on daily care (diet and exercise, etc.) to improving patient compliance

Even though there are many advantages pertaining to SNS-based community care websites, our study also has some limitations. First, the shared information may not be accurate or reliable. Because of registration by only e-mail, we cannot be sure that users are genuine DM patients. The effects caused by misleading information might lead to major problems. To minimize this problem, the medical staff continuously examined the “*My community-free board*” menu on the website. They also attempted to remove posts with serious ethical problems, unclear information, or suspected commercial intent. However, there were no problematic posts during the research period. The more people participate and post, the more effort the medical staff has to put into the website. However, if the community is large and activated, a self-purification process about misleading information is possible, and harmful health-related information would be filtered naturally. However, this does not mean that the self-purification would be flawless. Second, the small sample size is another limitation. This might be due to insufficient promotion for the website. We promoted it to patients who were prescribed insulin in our hospital and they joined in the website if they wanted to, so the number was too small.

5. Conclusion

On our website, patients suffering from the same disease, undergoing the same treatment, and with similar interests interact, share, and discuss health information and knowledge. Patients can compare their treatment with that of others and understand their disease status. As the patients share this sense of identity, they obtain new health information and learn about their diseases. Through this process, patients learn about their diseases not from the physicians but from among themselves. It is hoped that the patients would be motivated to participate actively by sharing. To achieve this, a new SNS system that can provide patients with health information through feedback and modeling and in which patients can participate and share information is needed. This is the expected role of “*I Love insulin*.”

Disclosure Statement

No competing financial interests exist.

Conflict of Interest

No conflicts of interest

Human Subjects Protections

Since the real names of the users are not used, the physicians are unable to search for the users. Our study did not use any private patient information or possess any possibility of additional physical risks to the patients involved. Owing to the anonymity of the data and the retrospective nature of the study, informed consent was not required.

Fig. 1 Web navigation

Fig. 2 Exemplary site of the screen a. "My Web-Chart" menu b. "My community-free board" menu

Fig. 3 Detailed list of contents by the "I love insulin" website user

Total	140 users
Male, n (%)	97 (69.3%)
Age, years	40.7 ± 11.7
> 65 years, n (%)	3 (2.1%)
HbA1c, %	8.1 ± 1.7
BMI (kg/m ²)	27.4 ± 11.8
Type of DM	
Type 1 DM	23 (16.4%)
Type 2 DM	85 (60.7%)
Not answer	12 (8.6%)
Unknown	20 (14.3%)
DM duration, years	8.5 ± 7.4
DM management	
OHA	43 (30.7%)
Only Insulin	68 (48.6%)
Insulin + OHA	29 (20.7%)

Table 1 Baseline characteristics of "I love insulin" website user

Table 2 Detailed list of contents by the "I love insulin" website users

	Question	Answer	Answer/Question ratio	p-value
Total number, n	77	57	0.741	
Diabetes itself, n (%)	18 (23.4%)	17 (29.8%)	0.944	Reference
Diet, n (%)	17 (22.1%)	1 (1.8%)	0.059	<0.001
One's glucose levels, n (%)	15 (19.5%)	1 (1.8%)	0.067	<0.001
Insulin regulation, n (%)	9 (11.7%)	2 (3.5%)	0.222	<0.001
Exercise, n (%)	6 (7.8%)	4 (7.0%)	0.667	NA
Web-page, n (%)	6 (7.8%)	1 (1.8%)	0.167	<0.001
No relationship to diabetes mellitus, n (%)	5 (6.5%)	31 (54.4%)		NA*
Oral medication, n (%)	1 (1.3%)	0 (0.0%)		1.000

*The frequency analysis did not include a case whose ratio was over 1.

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