

of IOL selection is crucial to post-operative quality of life. Patients should be informed of the benefits and risks, and also consider their personal values and preferences, to make an appropriate decision. Patient decision aids (PDA) are evidence-based tools designed to help patients make specific and deliberate choices among health-care options.² Presently, most PDA in use are paper-based and less approachable. Nonetheless, in the past decade, the practices of healthcare have moved toward an era of information technology. Therefore, computer-based decision aids are emerging in various fields of clinical practice.³ Although doctor-patient communications are essential in the shared decision-making process, computer-based decision aids may help patients to properly digest the vast amount of information before their discussion with doctors. The advantages of computer-based over paper-based decision aids are their ability to interact with users. Syrowatka et al had identified six important features of computer-based decision aids: content control, tailoring, patient narratives, explicit values clarification, feedback, and social support.⁴ In an attempt to apply these features in clinical practice, we developed and evaluated a computerized patient decision aid (cPDA) built on an interactive robot, to help patients in the selection of an appropriate IOL model before cataract surgery.

Materials and Methods

Development of the PDA

We searched for clinical researches published in English for IOL selections on PubMed between 2000 and 2019. The following keywords were used, “intraocular lens”, “monofocal”, “multifocal”, and “extended depth of focus”. The retrieved articles were reviewed for their level of evidence. The information regarding efficacy, safety, convenience, and cost were extracted from the reviewed articles.^{5–10} Important contents were adapted for the PDA, as recommended by the International Patient Decision Aids Standards (IPDAS) Collaboration.^{11,12} The PDA consisted of five parts. Part 1 is an introduction of the disease of cataract and its surgical management with IOL implantation. Part 2 lists all the IOL options and the strengths and weakness of each IOL model, including a tabular comparison of various options. Part 3 is comprised of a questionnaire to help patients rank their values and personal preferences of their visual needs in work and daily life. Part 4 is a small quiz to test the knowledge learned. Part 5 is a recommendation to the patients after considering their preferences and the knowledge they learned in the previous steps.

Computerize the PDA on a Robot

The cPDA was built on an interactive robot Zenbo (ASUSTek Computer Inc., New Taipei City, Taiwan). Zenbo is an Android-based robot equipped with a screen, speakers, camera, and multiple sensors to interact with users. The PDA were transformed into a user-machine dialogue with dialogue development environment (DDE) Editor and Zenbo APP Builder. The DDE Editor is a web-based interface to create and integrate conversational interfaces. The Zenbo APP Builder is a visual programming tool that allows developers to edit the codes by simply drag-and-drop-blocks, to control the flow and action on Zenbo. The interactive robot Zenbo can have dialogues with users through on-screen display and voice communication.

All five parts of the PDA were integrated into the cPDA. The cPDA first provides an introduction of the disease of cataract and its surgical management with IOL implantation. Then, it lists all the IOL options and the strengths and weakness of each IOL model, including a tabular comparison of various options, and will give a small quiz to test the knowledge learned. The quiz was in the form of a true or false test. If the patients failed the quiz, the cPDA would return to the prior sessions and then repeat the quiz. If the patients passed the quiz, the cPDA would proceed forward. After initial educational steps about relevant knowledge, the cPDA guides the patients through a series of questionnaires designed to help the patients assess their visual needs in detailed tasks, during sports, driving at night, desire to be glasses-free, as well as their financial concerns. The interactive cPDA provides specific questionnaires depending on the patient's choice of preference. In the end, the cPDA summarizes the preferences and recommends an IOL selection (Figure 1).

Pilot Testing of the cPDA

We conducted a pilot study for the helpfulness of the cPDA for decision-making in IOL selection before cataract surgery. The study was approved by the Institutional Review Board of Ditmanson Medical Foundation Chiayi Christian Hospital

Algorithm for Assessment of Visual Needs

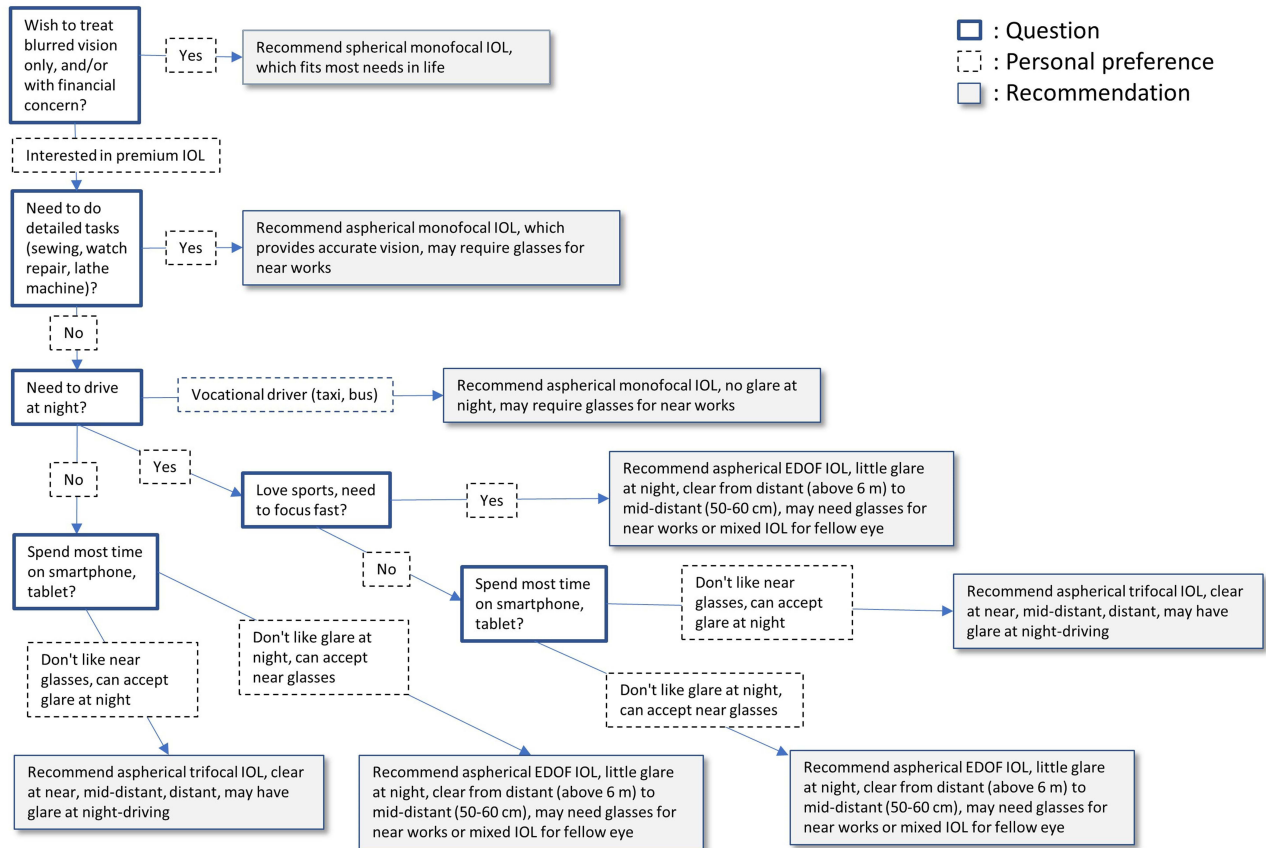


Figure 1 Algorithm for assessment of visual needs adopted by computerized patient decision aids.
Abbreviations: EDOF, extended depth-of-focus; IOL, intraocular lens.

(Permit No. 2020102) and was conducted in accordance with the Declaration of Helsinki. Patients and their families who were making the decision on IOL selection before cataract surgeries were eligible to participate in this study. Participants of different education levels were enrolled if they met the following criteria: (1) age of at least 20 years; (2) ability to speak and understand the Mandarin language; (3) no prior experience of decision-making in IOL selection. Informed consent was obtained from each participant in the study.

To test the helpfulness of the cPDA, we utilized two measurement tools, the Decision Self-Efficacy (DSE) scale^{13,14} and the Preparation for Decision Making (PrepDM) scale.¹⁵ Both scales and their Chinese versions were developed by the Ottawa Hospital Research Institute. The DSE scale consists of 11 items and the PrepDM scale consists of 10 items. Before the decision-making process, the participants complete the first half of the questionnaire for the pre-cPDA DSE scale. After receiving an introduction of basic operations, the participants went through the decision-making process with the aid of the cPDA (Figure 2). Each participant was tested individually in a separate room, to avoid interference. As the cPDA on robot Zenbo was interactive, the participants could order the robot to repeat certain steps of cPDA if they did not fully understand the information provided. After the process, the participants completed the second half of the questionnaire for the post-cPDA DSE scale and the PrepDM scale.

Results

A total of 50 participants (18 men and 32 women) were enrolled in the pilot test. The average age was 68.9 (range from 33 to 78) years. The patients with elementary education or below consisted of 44%, those with junior or senior high were 40%, and those of college or above education comprised 16% (Table 1).



Figure 2 A participant underwent the decision-making process using the computerized patient decision aids on an interactive robot. The robot was equipped with a touchscreen, speakers, camera, and multiple sensors to interact with users. In the photo, the screen is explaining how the intraocular lens can treat the cataract via audio and graphic expression.

Difference of DSE Scores Before and After the cPDA

The pre-cPDA and post-cPDA scores on DSE were computed as the sum of item scores divided by 11 and multiplied by 25. The results were stratified into different education levels (Table 2). The mean for pre-cPDA DSE score was 46.5 and the standard deviation was 13.6. Its distribution was skewed to the right. The post-cPDA mean DSE was 72.6 and the standard deviation was 12.8. Its distribution was slightly skewed to the left. The average gain score on DSE was 26.1 and the standard deviation was 8.0. The Shapiro–Wilk test for normality indicated that the gain score on DSE was normally distributed ($p=0.30$). The 95% confidence interval for the gain score on DSE ranged from 23.8 to 28.4. In summary, the gain score on DSE was statistically significant and the effect size was more than three.

Age, Sex, and Education Level in Relation to Gain Score on DSE and PrepDM

The PrepDM scores were computed as the sum of item scores divided by 10 and multiplied by 25. Multiple linear regression of the gain score on DSE and PrepDM on age, sex, and educational level is presented in Table 3. The patients with junior or senior high degrees had the highest gain score on DSE, followed by those with elementary school degrees or below, while the patients with college or above degrees had the lowest. The difference between the highest and lowest gain score was 7.4 and the associated 95% confidence interval was 0.3–14.6. Age and sex were not significant predictors

Table 1 Basic Information of Participants

Education Level	Male			Female		
	N	Age (Years) Mean (Range)	BVA Mean	N	Age (Years) Mean (Range)	BVA Mean
Elementary or below	5	82.0 (79–87)	0.29	17	70.7 (63–79)	0.45
Junior high	8	72.5 (63–80)	0.44	5	65.6 (61–69)	0.75
Senior high	2	68.5 (59–78)	0.67	5	66.0 (55–70)	0.8
College or above	3	66.7 (48–81)	0.7	5	51.8 (33–80)	0.63
Total	18	68.9 (48–87)	0.47	32	73.7 (33–80)	0.56

Abbreviations: BVA, visual acuity of the better eye in decimal scale; N, number of participants.

The cPDA used in the present study had the advantages of an initial interactive education stage, the ability to provide specific questionnaires depending on the patient's choice of preference, and to provide a summary of preferences and a recommendation of IOL selection at the end of the decision aid. The preliminary results showed the cPDA might help the self-efficacy of the patients in decision-making. There are still disadvantages of the present cPDA. Only one language was available. The limited items in choices for preference did not allow patients to fully express their values and preferences. The status of health literacy of patients was not evaluated before the decision aids.

The interpretation of the analysis results must consider the limitations of this study. Due to the lack of a control group, it was not clear how much the gain was actually due to the cPDA. The sample size was small and limited to one institution. Factors that might affect the patient's proficiency in using the cPDA, such as visual acuity, preferred language, cognitive ability, the presence of a doctor during the process, and other possible confounding factors were not studied. Whether the results could be replicated warrants further investigation. These limitations provide an indication for fields of further research.

Cataract is a disease of aging populations. Studies have shown that older patients tend to seek less information to make decisions and make decisions faster than younger individuals. Older patients have preferences for fewer choice options, face greater difficulties in understanding information about available options, and tend to disproportionately focus on emotional aspects when making decisions. Older patients tend to have variable preferences depending on their frailty, level of education, or cognitive and health status. Moreover, some, usually older and less educated, prefer to delegate decisions to their clinician.²² Most patients never knew about the complex knowledge of optics in the selection of IOL. A computer-based decision aid might help patients to assess their visual needs, clarify their values and preferences, and approach an optimal choice. However, it is the clinician's responsibility to accompany the patients through the whole process of decision-making, to maximize post-operative satisfaction in cataract surgeries.

Conclusions

An interactive cPDA may be a promising tool for complex decisions such as IOL selection before cataract surgeries. The preliminary results showed that education levels may be associated with the usefulness of cPDA on the preparedness and decision efficacy of the patients, although many confounding factors, including language preference, cognitive ability, and health literacy, need to be studied to support the clinical application of cPDA. This pilot study may provide a proof-of-concept of the feasibility of cPDA for patients who are making decisions of IOL selection before their cataract surgery.

Ethics Approval and Informed Consent

The study was approved by the Institutional Review Board of Ditmanson Medical Foundation Chiayi Christian Hospital (Permit No. 2020102) and was conducted in accordance with the Declaration of Helsinki. Informed consent was obtained from each participant in the study.

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Disclosure

The authors declare no conflicts of interest regarding the publication of this paper.

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