

Development of a Tool to Identify Patients' Preference for Vitamin K Antagonist or Direct **Oral Anticoagulant Therapy**

Shabnam Zolfaqhari, MSc¹ Job Harenberg, MD, PhD¹ Lutz Froelich, MD² Martin Wehling, MD, PhD¹ Christel Weiss, PhD³

¹Department of Clinical Pharmacology, Medical Faculty Mannheim, Heidelberg University, Mannheim, Germany

² Department Geriatric Psychiatry (CIMH), Medical Faculty Mannheim, Heidelberg University, Mannheim, Germany

³Department of Biometry and Statistics, Medical Faculty Mannheim, Heidelberg University, Mannheim, Germany

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Abstract

Keywords

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Address for correspondence lob Harenberg, MD, PhD, Department of Clinical Pharmacology, Medical Faculty Mannheim, Heidelberg University, Maybachstrasse 14, 68169 Mannheim, Germany (e-mail: job.harenberg@medma.uni-heidelberg.de).

Direct oral anticoagulants (DOACs) were developed for the treatment of thromboembolic diseases to overcome limitations of vitamin K antagonists (VKA). International guidelines on atrial fibrillation acknowledge patients' for antiembolic therapy with VKA or DOAC as relevant decision criteria. The objective assessment of patients' preference social interactions and psychological factors are hard to measure albeit representing important contributors. After a series of structured interviews and pilot studies assessing the preparedness to use DOAC as an anticoagulant and the motivation of patients to participate in clinical studies with DOAC, seven items were identified from several questionnaires by regression analysis. Those items were seen the best to describe the willingness to change anticoagulation from VKA to DOAC. By their use, we aim to develop a tool for the objective identification of the patients' preferences for VKA or for DOAC to increase adherence to therapy and to reduce anticoagulant undertreatment. German-speaking patients were asked to fill out a questionnaire consisting of biographic data and the seven selected items, and 180 patients completed the guestionnaire so far. Of these, 90 patients were on treatment with VKA (group 1), 57 patients changed anticoagulation from VKA to DOAC (group 2), 29 patients were DOAC naive patients (group 3), and 4 patients changed from DOAC to VKA (group 4). Overall, 94 patients received VKA, 29 patients received dabigatran, 50 patients received rivaroxaban, and 7 patients received apixaban as an anticoagulant. Eight patients anticoagulation were younger than 40 years, 35 patients were aged between 40 and 59 years, 53 preference patients were aged between 60 and 70 years, and 84 patients were aged older than adherence 70 years. Indication for anticoagulation were atrial fibrillation (n = 106), pulmonary embolism (n = 24), deep vein thrombosis (n = 40), artificial heart valve replacement (n = 8), or other diseases (n = 2). Based on the results of the analysis, a score will be dabigatran rivaroxaban suggested to identify the preference of patients for anticoagulation with VKA or DOAC. apixaban This tool may be useful for practitioners and health-care professionals to support patient atrial fibrillation adherence to therapy, and thereby increase treatment effectiveness.

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License terms (cc) (i) (=) (\$) Patients with nonvalvular atrial fibrillation (AF) are exposed to an increased risk for ischemic stroke and systemic embolism. Morbidity and mortality following stroke associated with AF are higher than in patients who suffer thromboembolic strokes in the absence of AF.^{1–3} Therefore, AF is one of the most frequent reasons for therapeutic oral anticoagulant therapy with vitamin K antagonists (VKAs) in everyday practice. For decades, VKA used to be the state-of-the-art anticoagulant to prevent thromboembolic events in patients with AF and additional risk factors.⁴ The major adverse effects are severe bleeding complications including intracranial hemorrhage.^{5,6} VKA usage requires frequent dose adjustments according to the international normalized ratio (INR) values between 2 and 3 to optimize the time in therapeutic range.⁷ Other downsides of VKA are interactions with food and many drugs, severe bleeding complications (intracranial and extracranial), and other severe adverse effects such as coumarin-induced hepatitis. The slow onset and offset of action of VKAs necessitates the overlapping administration of heparins and low-molecular-weight heparins during the induction of anticoagulation as well as during surgical interventions.⁸ Because of the limitations of therapy with VKA, patients are at risk for bleeding complications or for other reasons may not be treated with VKA or receive less effective medications, such as aspirin or even no prophylaxis.⁹

New anticoagulants (direct oral anticoagulants [DOAC]) were developed to overcome these drawbacks of conventional anticoagulant therapy and thereby help improve patient care. Advantages of novel synthetic DOACs inhibiting specifically only one activated coagulation factor include rapid onset and offset of action, lack of need for routine monitoring and dose adjustment, lack or infrequent food and drug interactions, and overall better efficacy and safety than the conventional anti-coagulants when used appropriately. The rapid onset and offset of action may eliminate the need of bridging with subcutaneous heparins as is required during anticoagulation with VKA.^{10–12}

For over 50 years of anticoagulation with VKA, patients did not have any choice for a different effective oral anticoagulant. This has now changed because of the development of DOAC, and now patients have the option to choose between VKA and DOAC. To qualify for this important decision-making process, patients and practitioners have to be well informed about advantages and limitations of available oral anticoagulants. Up to now, practitioners have largely acted as peers for patients which drug to choose for treatment.^{13–15} Preferences of patients for therapy with conventional VKA or for a DOAC may become increasingly important and this is mentioned in the guidelines of the American College of Cardiology Foundation (ACCF), American Heart Association (AHA), Heart Rhythm Society (HRS),¹⁶ European Society of Cardiology (ESC),¹⁷ and American College of Chest Physicians (ACCP).¹⁸ The understanding of the complex differences between oral anticoagulants by patients as laypersons is limited for many reasons, and seems similarly restricted as in patients with insulin-dependent diabetes.¹⁴ Some investigations have addressed this and correlated the patients' preferences with the willingness to pay for DOAC,¹⁹ the preparedness of

patients to use DOAC instead of VKA for anticoagulation,^{20,21} and the motivation of patients to participate in clinical studies with DOAC²² using questionnaires that were developed according to published techniques.²³ The patient preferences regarding treatment of human immunodeficiency virus (HIV)/AIDS identified a high impact of quality of life on patients' preferences for the selection of treatments was identified.²⁴ Furthermore, a tool was described to improve the adherence to HIV treatment based on psychological and environmental factors.²⁵ Involving patients in discussions about their treatment could lead to improved health through better adherence to chosen options, reduced practice variation about preference-sensitive options, and improved care in general.²⁶ So far, the willingness to change anticoagulation from VKA to dabigatran²¹ has been reported so far but not the preference of patients to choose between VKA and other DOAC. As a consequence, no tool is available to scientifically describe the preference.^{18,27,28} Earlier publications address the impact of patients' view on therapeutic options in anticoagulation in the pre-DOAC era or represent a personal view in the new era.²⁹⁻³¹ On this background, we aim to develop a tool to identify the preference of patients for a VKA or DOAC thereby improving patient's adherence to therapy as an essential prerequisite for success.

Development of the Questionnaire

Phase 1: Focus Groups for Concept Elicitation and Item Generation

The goal of this phase of development of the questionnaire was to elicit patients' experiences with oral anticoagulation for the treatment of AF and venous thromboembolism (VTE).²³ This phase was performed in a German focus group of patients on treatment with VKA using a structured discussion guide. Patients were recruited from the 4th Department of Medicine of the Medical Faculty Mannheim. The inclusion and exclusion criteria, ethics and informed consent procedure are described in the section "Study Design." Patients were invited to share their recollections, opinions, and wishes for anticoagulation, followed by discussions aimed at patients undergoing treatment with oral anticoagulation.

A set of items was developed that reflected the anticoagulation-related concerns of the patient. All major areas of comments that had been identified by patients were represented in the initial set of questions. A self-developed questionnaire was thus developed. It was then reviewed by an expert psychiatrist (L.F.) and modified accordingly. The questionnaire was divided into two modules: a biographic module and an anticoagulation module. The latter investigates the personal experience of patients with anticoagulation and wishes for an optimal anticoagulant drug.³²

In addition, patients' personality traits, fears related to a new medication, and self-assessment of the general health status were investigated by the validated personality inventory questionnaire Freiburger Persönlichkeitsinventar (FPI-R, Hogrefe-Verlag GmbH & Co.KG, Goettingen, 2001, Germany),³³ State-Trait Anxiety Inventory (STAI, Consulting Psychologists Press. Inc., Palo Alto, Calif., Beltz Test GmbH, Goettingen, Germany; 1970),^{34–37} and a 12-item short-form health survey questionnaire (SF-12; Hogrefe-Verlag GmbH & Co. KG Goettingen, Germany; 1998), summarizing two scales describing mental and physical well-being components.^{38–40}

Phase 2: Applicability and Acceptability of the Questionnaire

A total of 110 patients, on stable treatment with VKA for at least 3 months, completed the four questionnaires (FPI-R, STAI, SF-12, and the self-developed questionnaire).³² Patients replied to all questions indicating an adequate applicability and acceptability. Thus, rephrasing of the questions of our self-developed questionnaire was not required for further investigations.^{23,41}

Phase 3: Regression Analysis and Reduction of Items

In the meantime, DOACs were approved for prevention of embolism in nonvalvular atrial fibrillation. The wishes and fears for anticoagulation were discussed in this context with another focus group of patients in a structured interview under the guidance of an expert (J.H.). It is important to note that the price for DOACs is covered by health insurance companies in German-speaking countries. In this phase of development of the questionnaire, patients were investigated as to their disposition to change anticoagulant therapy from VKA to DOAC. Logistic regression analysis identified "extraversion" on the FPI-R being positively related to the willingness to change the medication. In addition, the following items of the self-developed questionnaire were identified: past thoughts regarding alternatives for anticoagulation, hope for a better quality of life, difficulty to adjust therapy based on the prothrombin time, lack of need for monitoring for dose adjustment of the anticoagulant drug, importance of the opinion of the practitioner on which anticoagulant to use, and skepticism to change drug for therapy.²⁰

Phase 4: Reliability of the Seven Items

To analyze the reliability of the results, 85 of 110 patients replied for a second time to the seven items. The receiver-

operative curve (ROC) confirmed the reliability of the seven items with a probability of 98%.²⁰ This high reliability supported the use of these seven items together with biographic data in patients' transition from VKA to DOAC (**-Table 1**).

Study Design

This is a prospective, controlled, multicenter, case-control study in patients with an indication for anticoagulant therapy. The Ethics Committee II of the Medical Faculty Mannheim, Heidelberg University approved the study protocol. For the paper version, patients had to give written informed consent. In the online version patients accepted participation by clicking the button "participate" implying to have read and accepted the informed consent in an accessible PDF document.

Patients: Four groups of patients are included in the study: patients on treatment with VKA (group 1), patients who switched anticoagulant therapy from VKA to DOAC (group 2), DOAC naive patients without personal experience with therapy of VKA (group 3), and patients who changed therapy from DOAC to VKA (group 4).

Inclusion criteria: The inclusion criteria were confirmed diagnosis of AF, deep vein thrombosis, pulmonary embolism or artificial heart valve replacement, oral anticoagulant therapy, aged 18 years and older, and the ability to understand German language.

Exclusion criteria: The exclusion criteria included severe diseases with impairment of cognitive function and age < 18 years.

Project overview: Patients undergoing treatment with VKA or DOAC can participate in the study. The projected inclusion period was until the end of 2013. The question-naire was divided into part one, referring to biographic data, and part two, referring to the seven questions (**-Table 1**). The aim of the project is to support a structured and justifiable decision on whether a patient prefers to continue with the current anticoagulant or to switch to the other group of anticoagulants based on the results of the seven items (**-Fig. 1**).

Part 1: Biographic data	Part 2: Items
 Gender Age Graduation level Diagnosis requiring anticoagulation treatment Anticoagulant currently being taken Duration of therapy with current anticoagulant Self-monitoring (if applicable) Reason for change of anticoagulant (if applicable) 	 Past thoughts regarding alternatives for anticoagulation Hope for a better quality of life Difficulty to adjust therapy based on the prothrombin time (INR) Lack of need for monitoring for dose adjustment of the anticoagulant drug Importance of the opinion of the practitioner on which anticoagulant to use Skepticism to change drugs for therapy Judgment of own personality trait for extraversion

Table 1 Parts 1 and 2 (7 items) of the questionnaire on the identification of the preference of patients for VKA or DOAC

Abbreviations: DOAC, direct oral anticoagulant; VKA, vitamin K antagonist.



Fig. 1 Outline of the study: structure of the questionnaire and decision tree to continue or to switch anticoagulant drug. DOAC, direct oral anticoagulants; VKA, vitamin K antagonist.

Methods

Generation, Documentation, and Processing of Data

Network

A network was organized for inclusion of patients into the project. A flyer was designed for distribution and publication in scientific and other journals. General practitioners, specialized hospital and outpatients care units, and blood coagulation centers agreed to participate. Advertisements were made at presentations on congresses, published in local and regional German newspapers in print, or on Internet forms and homepages of Gesellschaft für Thrombose und Haemostaseforschung with a link to www.blutverduennung.uni-hd.de.⁴²

Construction of the Questionnaire

The targets for appropriate construction of questions consist of (1) easy understanding, (2) simple feasibility, (3) shorttime solution, (4) low material and paper consumption, (5) easy readability, and (6) low frequency of random solutions.⁴³ For the answers to the seven items, we evaluated different scale formats in preliminary experiments (not shown) and finally chose a Likert scale for each of the items. Patients have to decide by agreeing or disagreeing with the statements of the items by choosing an integer between 0 and 10. This scale can be easily understood, offers graduated response choices depending on the intensity of agreement, and is readily interpretable.^{44,45} The time to answer to the whole question-naire including biographic data ranges from 5 to 10 minutes.

Processing of Data

The questionnaire on Internet was created using the Webbased software evaluation system (EvaSys, Electric Paper Evaluationssysteme GmbH, Lueneburg, Germany). EvaSys is an Internet technology-based system that can be used in particular scientific evaluations of educational events and seminars. The evaluation process with EvaSys is visualized in five successive phases. First, the design of the questionnaire is created. Second, the type of answers from various formats as single or multiple choice, open question, or scale is selected. The scales can be classified as nominal scale, ordinal scale, interval scale, and ratio scale (>Table 2). EvaSys generates a PDF file or an HTML file including patients' data and simple corresponding statistical measures such as frequencies or mean values. To ensure that only qualifying persons use the questionnaire, EvaSys individual transaction numbers (TANs) have to be created. On agreement of a person to participate in questionnaire, the TANs will be sent with a link from the homepage to the e-mail address of the participant. In the third phase, the data of the completed questionnaires are collected by EvaSys and automatically transferred into an Excel database spreadsheet (Microsoft, Redmond, WA). In the fourth phase, the raw data are exported for analysis in statistical programs such as SPSS (IBM Corporation, Armonk, NY), SAS (SAS Institute Inc., Cary, NC), or SPHINX (Sphinx Technology, Inc., Brooklyn, NY). The data analysis in our study will be performed with SAS statistical software release 9.3. Documented data in Excel fundamentally simplify the work for SAS programmers because it can import the data into SAS for evaluation. The fifth phase allows a quality management and process optimization of the system.

Statistical Analysis

Each question will be analyzed with a different statistical test depending on its special form. To compare the four groups regarding binary data (i.e., patients' gender or diseases), the Chi-square test or Fisher exact tests will be used, as

Scales	Advantages	Limitations	Examples
Nominal scale	Easy to understand and to determine	Not specific; binary data: high probability of randomly correct solutions	Gender
Ordinal scale	More graduations than nominal scale	May be misunderstood by responder; less precise than quantitative data	School grade, visual analog scale
Interval scale	Allows efficient analysis, differences between two subjects may be calculated	Requires validated techniques; ratios are not useful	Body temperature
Ratio scale	With absolute zero, only positive values; allows most efficient analysis, subjects may be compared by their ratios	Requires validated techniques	Age

Table 2 Description of the different forms of scales to quantify agreement or disagreement with a statement of a questionnaire



Fig. 2 Distribution of age of patients according to the gender included in the study (as of August 2013).

appropriate. For quantitative data, which are approximately normally distributed (i.e., patients' age), one-way ANOVA (analysis of variance) will be performed. The Kruskal–Wallis test will be applied for ordinally scaled (i.e., the seven items) or skewed data. Comparisons between the two groups will be done by the two sample *t*-test or Mann-Whitney *U* test. Furthermore, for each patient, a multiple logistic regression analysis model will be performed to assess patient's preference expressed by the probability to perform anticoagulation with VKA or DOAC. This has already been done in a previous study.²⁰

Preliminary Results

From February 1, 2013, to August 1, 2013, a total of 180 patients (109 males, age [mean \pm standard deviation or SD]: 66.9 \pm 10.9 y; 71 females, age [mean \pm SD]: 63.2 \pm 15.3 y) participated in the study and completed the questionnaire. The distribution of age in decades and according to gender is shown in **~Fig. 2**. A total of 60 patients have answered the questions on paper format and 120 patients on the online format. The school degree of education of patients is shown in **~Fig. 3**. Overall, 54 patients have elementary school certificates, 55 patients have intermediate school certificate, 24 patients have high school certificates, and 47 patients have university degrees. The indication for anticoagulation was



Fig. 3 Education level of the participants of the study (as of August 2013).



Fig. 4 Indication for anticoagulation of patients included into the study is given (as of August 2013).



Fig. 5 Distribution of anticoagulants used in the study (group 1, VKA; group 2, patients who changed from VKA to DOAC; group 3, DOAC naive; and group 4, patients who changed from DOAC to VKA). DOAC, direct oral anticoagulants; VKA, vitamin K antagonist.

mainly AF, followed by deep vein thrombosis, pulmonary embolism, and artificial heart valve replacement (**- Fig. 4**).

A total of 90 patients were on treatment with VKA (group 1), 57 patients changed anticoagulation from VKA to one of the DOAC (group 2), 29 patients were DOAC naive patients and placed straight onto DOAC (group 3), and 4 patients changed therapy from DOAC to VKA (group 4) (\sim Fig. 5). Patients of group 4 switched from DOAC to VKA due to fear of adverse effects on DOAC (n = 3) and one patient reported hair loss after 9 months of treatment. Of the 94 patients of group 1 and 4, 37 performed self-management of anticoagulation and 57 performed INR



Fig. 6 Distribution of patients on treatment with vitamin K antagonist (VKA) is depicted according to self-management of therapy or control of anticoagulation by the practitioner and according gender (as of August 2013).



Fig. 7 The use of the individual DOAC of patients according to the gender is shown. The doses of the DOAC are not identified because of low patient number (as of August 2013). DOAC, direct oral anticoagulants.

determinations at the general practitioner. Differences between genders cannot be identified (\succ Fig. 6). The distribution of the DOAC dabigatran (all doses), rivaroxaban (all doses), and apixaban (2×5 mg twice daily) for anticoagulation is given in \succ Fig. 7.

Discussion

The aim of the present, still ongoing study is to develop a tool for practitioners to identify the preference of patients for a VKA or a DOAC as anticoagulant to improve adherence of patients to therapy. Many studies reported that the quality of clinical communication is related to positive health outcomes. Communication between patient and physician helps improve and resolve the problem of choosing the type of anticoagulant therapy.⁴⁶ In particular lack of physicians' time may be one of many reasons why the patients' preferences for a specific anticoagulant are currently being overlooked. With this study, we aim to find a tool for the physician and the health-care system that helps patients to choose the preferred type of oral anticoagulant.

The proportion of patients using VKA as an anticoagulant due to a specific reason such as fear of increased bleeding risk with a DOAC may influence the patients' preference. If a patient would prefer DOAC based on the results of the questionnaire, the practitioner may decide in favor of DOAC for a plausible reason justifying the higher price. On the contrary, patients may prefer anticoagulation with a VKA based on the results of the questionnaire facilitating the choice of this type of the anticoagulant for an individual patient.

Currently the ACCF/AHA/HRS,¹⁶ ESC,¹⁷ and ACCP guidelines¹⁸ include the preferences of patients in their recommendations to choose VKA or DOAC for oral anticoagulation of patients with AF. Several reports are available on how to improve the adherence of a patient to anticoagulant therapy in patients with AF. It was recently reported if physicians' and patients' perspectives can be identified by answering several questions: "Do patients' and clinicians preferences explain the utilization of warfarin to prevent strokes associated with AF?," "To what extent do patients' and clinicians' treatment preferences differ?," and "What factors explain any differences that exist in treatment preferences between patients and clinicians?⁴⁷ Some guestionnaires were developed without using results of structured interviews with patients regarding the price patients would be willing to pay for the treatment with a DOAC.^{19,48} In this report, a questionnaire was developed using published methodologies based on structured interview with focus groups with the treatment in guestion.³² Thus, we have developed a set of approximately 28 questions related to therapy with VKA. In addition, standardized questionnaires regarding personality traits, anxiety, and self-assessment of health status were included. Logistic regression analysis identified seven items out of over 200 items regarding the willingness to switch anticoagulation from VKA to DOAC.²⁰ With this study, we want to identify the validity of seven items for patients who have really changed anticoagulation from VKA to DOAC. Thereby, the preference of patients for a specific type of an oral anticoagulant could be identified analyzing the differences between the four groups of patients.

An overview summarized 16 studies focused on patients with AF regarding the burden of antithrombotic treatment using different scales to identify the acceptance of thrombotic or bleeding risk on VKA.¹⁸ Around 20% were willing to accept 35 additional bleeds on VKA for 3% absolute risk reduction of stroke. For these 20%, the disutility associated with one stroke was equal to the disutility associated with 11 bleeding episodes.⁴⁷ Patients with AF placed more value on the avoidance of stroke and less value on the occurrence of bleeding in contrast to physicians who placed more value on the occurrence of bleeding during treatment with warfarin.¹³ This may relate to the concept of loss aversion, which refers to the tendency for individuals to prefer avoiding loss in favor of gaining benefits.⁴⁹

Methods used to convey information about risks and benefits of therapy may significantly affect patient-reported preferences.⁵⁰ Also, previous exposure to a therapy was associated with a preference for continuing the same treatment.^{51,52} Cognitive dissonance occurs when participants tend to collect additional information to ensure that it is consistent with their previous decision.⁵³ To reduce cognitive dissonance, participants may be inclined to continue their treatment, despite new information suggesting a suboptimal choice. Patients who do not want to believe that they have been taking the worse treatment may interpret the evidence presented in a way that it is consistent with their previous choice.¹⁸ These considerations can be translated to prefer anticoagulation with a VKA or a DOAC. This study should help overcome such cognitive dissonances for patients as well as practitioners.

Some limitations of the study should be considered. At present the investigations can be performed only with German-speaking persons. Translation to other languages encounters cross-cultural differences. International guide-lines were developed to adapt questionnaires to other countries and cultures.⁵⁴ The opinion of a patient about acceptable risks for a thromboembolic or a bleeding event and

economical aspects are not included in the questionnaire. In preliminary studies, these aspects were included and logistic regression analysis did not rank the results high enough to be included into the final present questionnaire.³² The higher daily costs for DOACs compared with warfarin are not included into the questionnaire because in Germany they are covered by the insurance companies. This is different in many other countries and has to be respected when the questionnaire will be implemented in other countries. Patients unwilling to participate in the study may prefer more frequently a conventional over a new DOAC. The noninclusion of these patients may cause a bias. Convincing as many patients as possible to take part in our study may counteract this. However, the problem of not participating is inherent in most studies and cannot be eliminated completely.

In conclusion, this investigation is undertaken to develop a tool for the health-care system to maximize the adherence of patients to anticoagulant therapy by identification of their preference for a specific type of an oral anticoagulant. Using this tool, patients will be actively involved in the decision to perform anticoagulation with VKA or DOAC. As a consequence of implementing the tool into patient care, undertreatment in anticoagulation should be reduced.

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Conflict of Interest

The authors do not declare any conflict of interest for the present investigation or preparation of the article.

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