Patients’ subjective concepts about primary healthcare utilisation: the study protocol of a qualitative comparative study between Norway and Germany

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ABSTRACT

Background: In Germany, utilisation of ambulatory healthcare services is high compared with other countries. While a study based on the process data of German statutory health insurances showed an average of 17.1 physician-patient-contacts per year, the comparable figure for Norway is about five. The usual models of healthcare utilisation, such as Rosenstock’s Health Belief Model and Andersen’s Behavioural Model, cannot explain these differences adequately. Organisational factors of the healthcare system, such as gatekeeping, do not explain the magnitude of the differences. Our hypothesis is that patients’ subjective concepts about primary healthcare utilisation play a major role in explaining different healthcare utilisation behaviour in different countries. Hence, the aim of this study is to explore these subjective concepts comparatively, between Germany and Norway.

Methods/design: With that aim in mind, we chose a comparative qualitative study design. In Norway and Germany, we are going to interview 20 patients each with qualitative episodic interviews. In addition, we are going to conduct participant observation in four German and four Norwegian primary care practices. The data will be analysed by thematic coding. Using selected categories, we are going to conduct comparative case and group analyses.

Ethics and dissemination: The study adheres to the Declaration of Helsinki. All interviewees will sign informed consent forms and all patients will be observed during consultation. Strict rules for data security will apply. Developed theory and policy implications are going to be disseminated by a workshop, presentations for experts and laypersons and publications.

BACKGROUND

The utilisation of ambulatory healthcare is comparatively high in Germany. The beneficiaries of statutory health insurances had an average of 17.1 contacts with physicians per person per year in 2007 in ambulatory care.1 This figure is considerably higher compared with those in other countries. The number of contacts for Norway, for example, amounts to 4.6 per person per year,2 a country which is comparable with Germany regarding mortality and morbidity among the population. The reasons for this significantly higher utilisation of ambulatory care in Germany compared with other countries such as Norway are largely unexplored. In this research project, utilisation of primary healthcare is understood as the number of contacts of a patient with a general practitioner (GP) per time unit.

Way back in 1966, Rosenstock posed the question ‘Why People Use Health Services’.3 He developed his so-called Health Belief Model on the utilisation of preventive and diagnostic healthcare provision. The Health Belief Model comprises two classes of variables: These are the readiness to get active and assumptions about the effectiveness of different measures.

The ‘Behavioural model on the utilization of healthcare provision’ published by Andersen for the first time in 1968 is also based on the idea that high-utilisation rates in healthcare are a positive phenomenon and will lead to better healthcare outcomes.4 Andersen distinguished predisposing factors such as demographic factors, social structure and health beliefs, enabling factors such as family, community and the perceived need for healthcare. Later on, Andersen extended his model with characteristics of the healthcare system (policy, resources, organisation) and fitted a feedback loop into his model. In Andersen’s model, health beliefs are defined as attitudes, values and knowledge about health and healthcare provision. However, quantitative studies which apply Andersen’s model rarely make use of the factor ‘health
beliefs’, which can mainly be attributed to difficulties in operationalisation. When it comes to the effect of attitudes, values and knowledge about healthcare provision between different countries within the framework of Andersen’s model, we do not know of any such study to the best of our knowledge.

On the basis of the models by Rosenstock and Andersen, numerous studies were conducted in order to get a better understanding of the predictors about individual utilisation behaviour. Exemplary studies focusing on the German healthcare model are the studies by Thode et al., Bergmann et al. and Kürschner et al. The results of these studies demonstrate the importance of age, morbidity and sex to the extent of utilisation of healthcare. Since there are no major dissimilarities regarding the factors age, morbidity and sex in the overall population in western countries such as Germany and Norway, these factors are unable to explain the vast differences in healthcare utilisation.

Another starting point for the explanation of utilisation behaviour relates to factors inherent to the healthcare system, specifically whether the GP functions as a gatekeeper to specialists. Garrido et al. examined in a systematic review the effect of gatekeeping on the utilisation of physicians working in ambulatory care. A tendency towards a reduced utilisation of specialists could be demonstrated, which to some extent was compensated by a higher recourse to GP. Hence, the effect of gatekeeping is not able to explain the magnitude of the differences found in utilisation of ambulatory care either.

Thus, we focus on health beliefs as defined in Andersen’s model, which have been widely neglected by scholars so far, as an important factor for different healthcare utilisation in different countries. A broader concept regarding health beliefs is subjective concepts. We view subjective concepts as underlying assumptions, values and knowledge with regard to a specific object. Hence, we define subjective concepts of the utilisation of ambulatory healthcare as the assumptions of patients, how and why they decide to make use of ambulatory healthcare given a specific health issue, as well as the underlying attitudes, values and knowledge leading to this decision. In turn, we see these subjective concepts as influenced by society. This definition of subjective concepts is guided by subjective layman theories, which contain an implicit structure of reasoning as described by Groeben et al. The second base is the ‘Theory of social representation’, which describes the societal contingency of subjective concepts.

As individual concepts of patients are (also) influenced by society, differing subjective concepts can be expected in different societies. Subjective concepts can therefore potentially explain differences in utilisation in several countries. Figure 1 depicts the theoretical framework of the study.

As a country of comparison for the concepts of German patients, Norway is well suited, because of its state-run healthcare system contrasting with Germany’s cooperative one with mainly self-employed GP in both systems. The cultural differences between Germany and Norway are small in scale, allowing for a comparison. Life expectancy in Norway can be compared with the one in Germany, whereas the utilisation of ambulatory care differs widely. With regard to experiences of physicians who have worked in both systems, as well as differences in medical guidelines, it can be presumed that Norwegian and German patients hold different views on the utilisation of ambulatory care.

Aims

Our aim is to research the patients’ subjective concepts which determine the different healthcare utilisation behaviour in Germany and Norway. Thus, our research question is:

How do adult patients’ subjective concepts about primary healthcare utilization differ between Germany and Norway?

Methods

Our methodological starting point is the patient’s own experiences and frequent reasons for encounter. The research design is qualitative: patients in Germany and Norway are going to be interviewed. In addition, we are going to conduct participant observation in primary healthcare practices. The data from both methodological approaches are going to be triangulated. We are going to compare cases and groups based on thematic coding.

Methodological approach

In order to conduct research on cultural differences, several methodological approaches are available: for example, participant observation is widely common in ethnography, and the episodic interview is used to compare groups by thematic coding. The underlying theoretical assumption in both approaches is that individual perceptions and concepts are at least partially socially constructed.
In this project, both methodological approaches shall be applied and triangulated. Figure 2 gives an overview of the project’s theoretical framework, completed by the respective data collection instruments: both episodic interview and participant observation apply different methodological approaches to research the subjective concepts of patients and the resulting behaviour. The usage of both methods and their triangulation shall lead to broad, rich and valid results.

**Data collection methods**

In many studies, qualitative interviews have been employed to research subjective concepts, since qualitative interviews provide access to individual cognitions and their world of experiences. In this context, Flick has developed the method of the ‘episodic interview’ specifically designed to compare the subjective concepts of specific groups. In episodic interviews, questions posed in order to evoke narrations are combined with questions aiming at argumentative-theoretical answers. The topics and the narrative stimuli will be covered by an interview guideline. The guideline does not imply a certain order of the respective questions, but rather serves as an aid to memory. The aim is to get into a conversation with the interviewee, which resembles a conversation, under which course the respective parts on the guideline will be touched upon. In this study, the guideline has already been developed (cf. box 1).

Participant observation as a method relies on being present in the field, that is, in the setting, among the individuals and surroundings to be researched. Next to the actors’ perceptions in the field, the action based on them is on the focus of this particular method. Thus, we intend to observe how patients utilising primary healthcare express their reasons for utilisation and assumptions of care in the GP’s practice. This can, for example, be at the reception desk, in the waiting area, during the consultation with the GP, in the interaction with other patients, the staff or the GP. The projected time in each GP’s practice is 1 week. The researcher attempts to get into a conversation with the actors involved, in order to explore their underlying point of view for their actions. A regular retreat of the researcher in order to note down the observations is a characteristic of the method and secures the quality of the data. In order to improve the quality and comparability of the participant observation, an observation guideline will be used, which lists the general framework and objectives for the observation as well as the notes. This guideline has already been developed (cf. box 2).

**Sampling**

Concerning the participant observation, we base our sample on a 2×2 design: Of the overall eight observations, lasting for 1 week each, half will be conducted in Germany, and the other half in Norway. In order to account for the potential differences in the availability of services and the socioeconomic standing of different types of municipalities, it will be differentiated between a rather urban setting and a more rural area in each country (cf. Refs. ). That yields the distribution of participant observations as summarised in figure 3. The 81-week participant observations facilitate a very broad design accounting for differences, while, at the same time, keeping the volume of qualitative data material within manageable limits.

The number of interviewees also aims at reaching a relevant amount of comparison with a contentwise saturation, which at the same time still allows for a deeper analysis of the collected data. Therefore, 40 patients shall be interviewed, half of them in Germany, and the other half in Norway. Figure 3 provides an overview of the sampling procedure.

Contact with potential participants in the study shall be established through the eight cooperating GPs’ practices. All patients visiting on any of two certain days will consecutively receive an information sheet asking for the declaration of consent to be contacted by the research team and a short survey on age, sex, subjective
morbidity and the number of contacts with a physician during the last 3 months. Out of this range of potential interviewees, the actual participants will be picked in an iterative process. Therefore, interviewees will be chosen evenly from both sexes and different age groups. In order to arrive at comparable results, German interviewees shall be matched with the Norwegian ones with regard to age, sex, subjective morbidity and healthcare utilisation. The German interviewees will be chosen according to the Norwegian participants. For matching a point, the system evaluates the match with a score of 5 points as best matching and 4 or 4.5 points as good matching. Table 1 displays the matching criteria. The quartiles concerning subjective chronic diseases and visits to a physician are calculated separately in each country for urban and rural areas.

After conducting half of the interviews and a first rough analysis, it will be decided whether or not patients generally not visiting any GP or specialist are sufficiently represented in the sample. It might be necessary to access them via a different sampling procedure in the second round of interviews.

Data analysis

The analysis of the qualitative data material is based on the concept of Grounded Theory.23 The basic underlying idea of Grounded Theory is to develop a specific, short-range or middle-range ‘theory’ in a stepwise process of analysis, that is, ‘grounded’ in the data. Theoretical coding is at the heart of the analytical process, for which codes are assigned to certain text segments. What is different from many other coding schemes is that the codes are not set beforehand but emerge during the process. Data collection and analysis occur step by step, not sequentially, but parallel with each other.

Based on theoretical coding, thematic coding was developed by Flick.24 An essential dissimilarity between the two of them is that the latter first takes individual cases into account and, second, facilitates comparisons between groups. While theoretical coding dissolves the single cases into the overall data material, the relation to the individual case is preserved in thematic coding. The reference to the case is ensured through a description of each individual case. In addition, it facilitates the development of typologies for specific aspects of the researched topic based on the respective cases. The project will make use of the aforementioned possibility of comparison of groups, where interviewees in Norway
and interviewees in Germany are going to comprise the two main groups.

The interviews will be transcribed word for word, after which a rough case description will be developed for each case. To begin with, when it comes to the analysis, two interviews from Norway and the same number from Germany will be chosen according to the method of maximising the contrast between the respective cases.  

The chosen interviews will be coded line by line, that is, an ad hoc code will be assigned to each line. The codes themselves are not predefined. Based on these line by line codes of the four interview transcripts, a thematic structure will be developed. A thematic structure is a system of categories, in which each code can be associated with a category. As a last step, all interviews will be coded subsequently within this thematic structure, which however will be adapted over the course of coding so that new categories can be added or sorted differently. Parallel to coding, the case description will be developed further and refined. The same applies to sequences of the notes taken during the observation.

Line-by-line coding is performed by the first and the second authors. They compare their line-by-line codes and discuss the differences and similarities. Coding is performed by the first and the second authors. The coding is reviewed in sessions by the project team. During the process of data analysis, we are going to write memos on the cultural and professional differences emerging. These memos and protocols of sessions of data analysis are collected and seen as part of the interpretation process and quality management.

Based on the thematic categories and the assigned codes and text segments associated with them, a thorough analysis of selected topics can be conducted. With the aid of comparisons between cases and groups, subjective concepts of patients in Norway and Germany can be compared and typologies developed.

**Ethical considerations**

The study adheres to the Declaration of Helsinki. All interviewees sign an informed consent as well as all the patients observed during consultation. Strict rules for data security apply: Addresses and contact information are stored physically separate from content data. Any contact information is destroyed as early as possible, for example, after the interview; only anonymised data are used for analysis. The study has been approved by the local ethics committees in Magdeburg/Germany and Bergen/Norway.

**Dissemination**

Developed theory and policy implications are going to be disseminated through a workshop for researchers and policymakers. The results are going to be presented at several congresses and conferences, and we are going to publish the results in peer-reviewed journals. In addition, we are going to present our results to laypersons by public talks.
DISCUSSION

Conducting a comparative qualitative study creates several difficulties which have to be addressed when planning such a study. An important problem in comparative studies is language. Qualitative interviews, participant observation and data analysis are highly language dependent. Thus, native speakers taking part in the study and having good knowledge of both languages of the core personnel are important for conducting a valid study. An important question is when in the process of analysis to translate the interviews. We decided to code the interviews in the original language and to translate the sections needed for publication in the last step. In our research group, three researchers and one student assistant are able to understand at least written German and at least written Norwegian. This seems fairly convenient to avoid language difficulties; however, it partly excludes those researchers who are not able to read in both languages.

Another difficulty is the funding of a comparative study. National grant agencies usually finance personnel in their own country only. EU funds are usually announced for specified topics and only for projects involving more than two countries. The organisation of data privacy and ethical reviews are different between different countries: for example, in Norway, the emphasis is on the privacy of the physician–patient encounter, while the emphasis in Germany is on data processing and use of personal data. Oeye et al.²⁷ have already described the difficulties in obtaining ethical approval for participant observation in the medical field in Norway.

CONCLUSION

In health services research, it is important not only to state the quantitative differences between countries, but also to look for the reasons for these differences. Qualitative methods are available to compare the subjective and social factors potentially involved in these differences or to explore the possible reasons for the differences. However, comparative qualitative methods are challenging because of the methodological and organisational difficulties occurring.

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