Qualitative Research in Nephrology: An Introduction to Methods and Critical Appraisal

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Introduction

Identifying and implementing patient priorities, values and goals in research, practice and policy is widely advocated to improve patient-centered care and outcomes. Qualitative research is a broad term for various approaches of inquiry that generate insights about peoples' perspectives, beliefs and attitudes, which can drive or explain behaviors, decision-making and outcomes. Such insights can inform strategies and interventions to address barriers and challenges with regards to clinician-patient communication, shared decision-making, education, delivery of care and resource allocation.

Qualitative methods may yield answers to important questions in nephrology and transplantation. For example, exploring patient attitudes and preferences towards vascular access in hemodialysis can help to address barriers to achieve timely creation of permanent vascular access, which is associated with reduced mortality, infection and healthcare costs. Furthermore, to address the gap between the need for and supply of transplantable organs, qualitative methods are used to describe community attitudes and beliefs towards deceased organ donation. This can assist in the development of education strategies, and policy change to increase deceased organ donation rates.

In recent years, there has been a substantial increase in the number of qualitative research articles published in medical journals, including in nephrology. Qualitative research conducted in a rigorous, systematic and transparent manner yields comprehensive, and compelling insights to inform practice and policy. This article provides an overview of qualitative research methods and a framework to appraise qualitative studies.

Overview of Qualitative Methods and Methodologies:

Overview
Qualitative research encompasses a variety of different methodologies and methods, these are outlined in Figure 1. Methodology refers to the theory or framework (e.g. grounded theory, phenomenology or ethnography) that researchers may use, which can inform the choice of methods (e.g. participant selection, data collection and data analysis). Usually, qualitative studies will involve an inductive approach, allowing the theories or concepts emerging from the data to generate hypotheses. This differs from the deductive methods used more often in quantitative research which centers around testing a pre-determined hypothesis. Qualitative research adopts an iterative approach, whereby researchers continually consult collected data throughout all stages of the study to inform their next steps. This may occur at the level of study design, such as revising the research questions or at the level of data collection where adjustments are made to the interview questions to clarify meaning and formulate concepts as new insights emerge. Qualitative research methods can also be combined with quantitative methods in a mixed-methods study. This complementary approach utilizes qualitative methods to explore quantitative findings to generate more complete data.

**Participant Selection**

Sampling in qualitative research seeks to obtain rich and diverse perspectives and meaning relevant to the research question. A purposive sampling strategy involves selection of research participants with particular characteristics of relevance such as ethnicity or disease group. Such sampling aims to include a range of demographic, clinical characteristics, experiences and backgrounds. This approach focuses on involving “information-rich” participants to obtain broad insights relevant to the research question.
Convenience sampling involves a more opportunistic approach whereby the participants selected are easily accessible. The snowball approach involves the identification of subsequent respondents by asking participants to identify individuals who can give important and relevant insights on the research topic. Theoretical sampling is used in grounded theory where participants are selected specifically to test a theory as it emerges during the data analysis.

**Data Collection**

Qualitative studies may use one or more of the following methods of data collection: unstructured or semi-structured interviews, focus groups, observations and document analysis. Semi-structured interviews are the most frequently used in health research and generally include 20-50 participants. Question guides are developed following literature and expert review and cover topics relevant to the aims and scope of the study. Open-ended questions prompt and encourage participants to elaborate on their responses. Such question guides may be pilot tested to ensure comprehension and clarity among a sample of study participants and further refined during data collection to address newly emergent topics and to capture the depth and breadth of data. Unstructured interviews are in-depth and take a more narrative approach, inviting participants to share their experiences with occasional prompts. Focus groups capitalize on group dynamics and encourage active participation and interaction among participants. Facilitated discussions encourage participants to share their views through shared experiences and explore similarities and differences of opinion. Focus group studies typically report three or more groups each with 6-8 participants to optimize participant interaction.

Observation studies the events and actions within a particular social or historical context. Observation can be used with other methods such as interviews, to obtain broader more comprehensive perspectives. Document analysis involves identification, organization, evaluation and
synthesis of documentary data studies (e.g. newspapers, social media content, speech transcripts and published policies).

**Data Analysis**

Qualitative data analysis involves the identification of patterns in the data to develop a descriptive and analytical framework. Data analysis is a systematic, interpretive and iterative process which involves reading the data, identifying/coding phrases, items and events and comparing codes throughout the data set. On completion of initial coding, the investigators group concepts into themes and identify patterns and relationships among themes. This iterative process aims to capture all concepts related to the phenomenon being investigated. It is important to acknowledge that the researchers’ background, experiences and values shape the process of data analysis and interpretation. Thus, the following strategies are implemented aiming to enhance the authenticity of the findings; member checking (seeking feedback from participants on the preliminary findings), investigator triangulation (involving multiple researchers who independently review the data to contribute to analysis) and reflexivity (acknowledging and examining researchers’ own biases that may affect the research process).²,⁵

**Reporting of Qualitative Research:**

Several guidelines are available for reporting and appraising qualitative research. We recommend the use of the Consolidated Criteria for Reporting Qualitative Research (COREQ)⁵ as a reporting guideline for interview and focus groups studies. The COREQ checklist was developed to promote complete and transparent reporting on aspects of qualitative studies (including details of the research team, study context, methods, findings, data analysis and interpretation).
Appraisal of Qualitative Research:

Qualitative research should generate comprehensive and trustworthy insights, whereby the research process is documented in a way that allows for auditing. As such, steps in the research process can be retraced to lead to certain interpretation or theory to ascertain that no alternatives were left unexamined.

The following constructs proposed by Lincoln and Guba can guide the appraisal of qualitative research: Credibility (Can the findings be trusted?), Dependability (Is the process logical and auditable?), Transferability (Are the findings relevant to other contexts and settings?) and Confirmability (Are the findings and interpretations linked to the data?). Table 1 provides a guide for assessing qualitative research according to these four constructs.

Credibility

Credibility refers to the extent the research offers comprehensive, trustworthy, and sensible explanations based on the data. Strategies to optimize credibility include providing a description of the research team, their roles, the study setting, the question guide and findings. This can assist in appraising whether factors individually or in combination influenced participant responses or investigator interpretations. The question guide should be relevant to the research question and designed to facilitate in-depth discussion. The study context and findings should be described in detail using “thick description”. This approach provides comprehensive contextual information of the research setting to enable readers to assess whether the results capture the depth and scope of the data. Member-checking and investigator triangulation ensures that analysis captures the range and depth of data.

Dependability
The concept of ‘reliability’ is not applicable to qualitative research methods, given the interpretive approach the findings are not able to be replicated or reproduced. Dependability, moreover, determines whether the research process is logical and auditable. In order to demonstrate that the study was conducted using a rigorous and systematic approach, it is important to clarify how the data was collected and analyzed in order to demonstrate a coherent link between the findings, methods and methodology. Use of audio or video recording, data transcription and coding software enables transparency and auditability of the research process.

**Transferability**

Whilst generalizability of results and external validity is an important criterion in quantitative research, this is not feasible in qualitative research since these studies necessitate small scale, in-depth conceptualized information. Transferability is the extent to which the concept and theories are relevant to other settings. The study should provide a detailed description of the study setting, participant characteristics, and healthcare framework. Furthermore, discussion of the extent the study findings resonate with published literature and how such findings advance theoretical understanding, can assist readers to determine the transferability to their own context.¹

**Confirmability**

Confirmability is defined as the extent to which the study findings reflect the participant’s perspectives, without being influenced or biased by the researcher’s predetermined assumptions or agenda. This can be achieved by researcher triangulation, member-checking, using quotations to link the findings to raw data and researcher reflexivity.¹
Examples of Qualitative Research in Nephrology:

Timely Creation of Permanent Vascular Hemodialysis Access

It is recognized that timely creation of permanent dialysis access is important to improve health outcomes in those transitioning to hemodialysis. However, the rates of patients starting dialysis with permanent access remain suboptimal. Richard et al.\(^7\) sought to examine the experiences of patients living with and caring for arteriovenous fistulas. An ethnographic study was conducted, which included semi-structured interviews of 14 purposely selected participants as well as extensive field work, and observations during vascular access and dialysis. Transcripts from audio-recordings and fieldnotes were analyzed through thematic synthesis. The overarching theme was vulnerability based on dependence on the healthcare system and the integrity of their vascular access for survival. Patients’ response to their vulnerability was heightened bodily awareness, vigilance and mistrust which led to stigma relating to their vascular access. The insights generated through researcher emersion into the culture of hemodialysis can inform future research and interventions to address these feelings of vulnerability and improve patient experience.

Adolescent Kidney Transplant Recipients Perspectives on Medication Adherence

The period of adolescence for transplant recipients is particularly challenging characterized by increased risk of nonadherence to treatment, loss to follow-up, and graft rejection leading to graft loss. Harwood et al.\(^8\) sought to explore the lived experiences of adolescent transplant recipients with respect to medication adherence. This phenomenological study conveniently sampled adolescent transplant recipients to participate in semi-structured interviews. Data analysis yielded an overarching theme; weighing risks and taking chances. Adolescents’ perceptions of the risks involved in non-adherence weighed heavily in their decision making. Modifications to their regimen were driven by protecting “normality”, peer reactions, and their self-concept and self-image. Such insights
into the lived experiences of adolescent transplant recipients is key to overcoming the challenges faced in practice.

**Shared Decision-Making in Selecting Dialysis Modality**

Morton et al. sought to identify, and rank characteristics of dialysis deemed most important to patients approaching dialysis and their caregivers. Thirty-four participants, including pre-dialysis, dialysis patients and caregivers were purposely sampled to partake in focus group discussions. A mixed-methods approach was adopted which included a nominal-group technique to obtain patient ranking of dialysis characteristics and qualitative thematic analysis to understand the reasons underpinning these rankings. Overall, patients and caregivers most value treatment which enhances survival, that can be undertaken at home. These findings can inform future work to enhance patient education and shared decision-making to improve patient experience and outcomes.

**Conclusions:**

The widespread shift towards improving patient-centered care and policy has highlighted the value of qualitative research in informing clinical practice and policy to improve health outcomes. Qualitative research methods elicit patients’ values, and attitudes that explain their health-related decisions and behaviors. However, qualitative research is time and resource intensive and is context-dependent so the findings may not be transferable to all populations and settings. As with all types of research, qualitative studies must be conducted using a rigorous approach with transparent and comprehensive reporting. Authors can aim to improve rigor by following well established principles and by demonstrating the key constructs of credibility, confirmability, dependability and transferability. Further contributions of qualitative research are needed to provide a broader evidence base to advance patient-centeredness in nephrology practice and policy, overall improving outcomes that are important to patients. Examples of topics that can be addressed with qualitative
research methods may include: the challenges of transplant recipients and caregivers during the COVID-19 pandemic, attitudes to telemedicine services for long-term follow up in transplantation, and shared decision-making in end-of-life care.

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References:


### Table 1: Proposed framework to guide appraisal of qualitative research

<table>
<thead>
<tr>
<th>Key Constructs</th>
<th>Definition</th>
<th>Strategies to satisfy criteria</th>
<th>Description</th>
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<tbody>
<tr>
<td><strong>1. Credibility</strong></td>
<td>Extent the research offers well founded, reliable and sensible explanations based on evidence obtained</td>
<td>Appropriate question guide</td>
<td>Relevant to research topic and phrased to encourage open in-depth discussion</td>
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<td></td>
<td>Data triangulation</td>
<td>Multiple sources of information and methods to produce a more comprehensive set of findings</td>
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<tr>
<td></td>
<td>Researcher triangulation</td>
<td>Multiple investigators from different disciplines with required knowledge and research skills to conduct the research</td>
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<tr>
<td></td>
<td>Purposive sampling</td>
<td>Recruit participants who can provide diverse and comprehensive information relevant to the research question</td>
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<tr>
<td></td>
<td>Theoretical saturation</td>
<td>Recruitment ceases when little or no new themes emerge from data already collected</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Thick description of data</td>
<td>Study context and findings discussed in detail</td>
<td></td>
</tr>
<tr>
<td><strong>2. Dependability</strong></td>
<td>Research conducted in a logical manner which can be audited.</td>
<td>Recording data</td>
<td>Audio or video recording to capture all data. Field notes capture contextual details and non-verbal communication</td>
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<tr>
<td></td>
<td>Verbatim transcription</td>
<td>Verbal data transcribed verbatim</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Computer software</td>
<td>Used to assist with storage, coding and retrieval of data</td>
<td></td>
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<tr>
<td><strong>3. Transferability</strong></td>
<td>Fit within contexts outside the study situation</td>
<td>Detailed description of research context</td>
<td>Detailed description of study setting and participant characteristics</td>
</tr>
<tr>
<td></td>
<td>Theoretical context</td>
<td>Findings are positioned within current theoretical or conceptual frameworks</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Comparison with other studies</td>
<td>Findings are within the context of other research conducted in different healthcare settings</td>
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<tr>
<td><strong>4. Confirmability</strong></td>
<td>Extent to which biases, motivations, interests or perspectives of the inquirer influence interpretations</td>
<td>Researcher triangulation</td>
<td>Multiple investigators involved in data analysis, ensuring the coding and analytical framework captures all primary data</td>
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<td></td>
<td>Member-checking</td>
<td>Participants provide feedback on preliminary findings which is incorporated into analysis</td>
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<tr>
<td></td>
<td>Inclusion of raw data</td>
<td>Selected quotations or other raw data inserted to illustrate findings or themes provided</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Researcher reflexivity</td>
<td>Researchers’ recognition of their own biases that may have influenced decisions made during the study</td>
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</table>
Figure 1. Common methodologies and methods in qualitative research

<table>
<thead>
<tr>
<th>METHODOLOGY &amp; EXAMPLES</th>
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<tbody>
<tr>
<td><strong>GROUNDED THEORY</strong></td>
</tr>
<tr>
<td>Use of systematic and iterative data collection and analysis to build theories about social phenomena</td>
</tr>
<tr>
<td>Community attitudes to deceased organ donation</td>
</tr>
<tr>
<td>Focus group study</td>
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</table>

| **PHENOMENOLOGY**       |
| Theoretical framework exploring how individuals make sense of the world providing insights of their “lived experiences” |
| Adolescents’ experiences of regimens after kidney transplantation | Harwood et al.* |
| Semi-structured and in-depth interviews |
| Medications made to the post-transplant regimen by adolescents were influenced by its impact on daily life and self-image |

| **ETHNOGRAPHY**         |
| Process to describe and interpret behaviors of social groups and cultures through observational and interview data |
| Barriers to nursing provision of patient-centered care in hemodialysis units | Tranter et al.* |
| Observations, interviews and case note reviews |
| Increasing demands of the unit led to focus on the dialysis machine and treatment thus neglecting the holistic care needs of patients |

| **NARRATIVE INQUIRY**   |
| Process of cross-disciplinary examination of stories of life experiences in the form of narrative interviews, diaries or biographies |
| Perceptions regarding death of individuals with chronic kidney disease and their families | Molzahn et al.* |
| In-depth interviews |
| Patients’ and families’ perspectives on end-of-life mostly align however, discussions with clinicians are much needed to elucidate preferences in care and planning |

| **CASE STUDIES**         |
| In-depth analysis of one or several individual cases with focus on circumstances, complexities and dynamics |
| Information needs of older people who choose supportive care over dialysis | Moustakas et al.* |
| Semi-structured interviews and case note review |
| Information regarding supportive care is infrequently delivered. As such older patients have little knowledge of health services to support them |

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**PARTICIPANT SELECTION**

**PURPOSIVE**
Selection of diverse research participants relevant to the relevant to the research question

**CONVENIENCE**
Selection of easily accessible and willing participants

**THEORETICAL**
Used in grounded theory where participants selected to test theory emerging in concurrent analysis

**SNOWBALLING**
Participants identified by study respondents, used to identify those in hard-to-reach populations

**DATA COLLECTION**

**INTERVIEWS**
Engaging in in-depth discussion to gain insights into individual perspectives

**FOCUS GROUPS**
Facilitated group discussion, capitalizing on group dynamics to generate data

**DOCUMENT ANALYSIS**
Identification, evaluation and synthesis of everyday media, health service policy, diary entries, or educational materials

**OBSERVATIONS**
Field notes collected of observations, videos or photographs

**DATA ANALYSIS**

**THEMATIC**
Derive and identify concepts and themes inductively from the data

**NARRATIVE**
Explores how people make and use stories to interpret experiences

**GROUNDED THEORY ANALYSIS**
Iterative data analysis in conjunction with data collection building theories grounded in the data

**CONTENT**
Deductive method of identifying codes prior to searching for them in data

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