Telemedicine in primary healthcare for the quality of care in times of COVID-19: a scoping review protocol

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ABSTRACT

Introduction Telemedicine gained strength in primary healthcare (PHC) during the COVID-19 pandemic. Thus, there is a need to know its scope, technologies used and impacts on people’s health. This study will map telemedicine use in PHC around the world and its impacts on quality of care in the context of the COVID-19 pandemic.

Methods This is a scoping review protocol developed according to Arksey and O’Malley and Levac et al, based on the Joanna Briggs Institute manual, and guided by the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Extension for Scoping Reviews (PRISMA-ScR). The records will be mapped in the following multidisciplinary health sciences databases: Virtual Health Library, PubMed, Scopus, Web of Science, CINAHL and Embase. Searches will also be conducted on Google Scholar, preprint repositories and specific COVID-19 databases (grey literature). Quantitative data will be analysed using descriptive statistics, while thematic analysis will be performed for qualitative data. Preliminary findings will be presented to stakeholders to identify missing studies and develop effective dissemination strategies.

Ethics and dissemination Results will be disseminated through publication in an open access scientific journal, scientific events, and academic and community newspapers. Ethical approval was obtained due to stakeholder consultation, but will not involve the direct participation of patients. Link to the protocol record in the Open Science Framework (OSF) (osf.io/q94en).

INTRODUCTION

COVID-19,1 caused by the SARS-CoV-2,2 was first reported in December 2019 in China. This disease triggered a public health emergency,3 leading the WHO to declare a pandemic situation.4 On 15 March 2021, the number of confirmed cases and deaths worldwide surpassed 119 and 2.6 million, respectively.5 Although the United Nations, WHO and Pan American Health Organization are working hard to obtain more information to combat the disease, they are still limited, and a greater level of knowledge is needed in all scientific areas.6

Following WHO recommendations, several countries have implemented quarantines,7 social distancing and mandatory home confinement.8 These measures have encouraged new technologies to respond to primary healthcare (PHC) demand. A tool that has been used is telemedicine, which has become a strategy to maintain assistance9–12 and guarantee admissibility and quality of care.13

The WHO adopted telemedicine or telehealth to define healthcare support using information and communication technologies (ICTs)14,15 in situations in which distance and/or geographical barriers hinder healthcare services.14 ICTs (ie, internet, cell phones, computers or satellite TV) make communication flows between healthcare professionals and patients more effective; thus, enabling real-time assistance.16 Also, ICTs offer relevant information related to asynchronous healthcare using electronic portals or other virtual resources, such as email, text messages and mobile phone applications.16

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Telemedicine use during the COVID-19 pandemic is occurring on a large scale and speed, becoming an efficient strategy to face PHC demand. This technology contributes to quality of healthcare services in PHC and strengthens monitoring, surveillance and detection of new COVID-19 cases. It also contributes to reducing patients' anxiety due to social isolation and maintaining contact between health professionals and patients with SARS-CoV-2; thus, allowing timely attention to most urgent cases and those with chronic diseases.

Regarding the healthcare team, telemedicine has shown a positive impact on workloads, with a reduction in burnout syndrome, which has been recurrent since the beginning of the pandemic. This technology has also contributed to reducing fear of virus exposure by professionals, collaborating with quality of care offered, facilitating coordination between primary and secondary healthcare levels, and maximising patient and family well-being in the context of the pandemic.

Quality of care has three dimensions: technical (action choice accuracy and production), interpersonal (social and psychological relationships between healthcare providers and users) and organisational (conditions in which services are offered, including continuity of care, coverage, action coordination, and service access and accessibility). In this study, technical dimension will represent the implementation of prevention and diagnosis measures and symptomatic treatment of mild COVID-19 cases in a timely manner (according to scientific protocols). Emotional support by healthcare professionals to users and/or family members in a situation of social isolation will represent the interpersonal dimension, while continuity of care for patients without COVID-19 and monitoring and follow-up of COVID-19 cases will be considered the organisational dimension.

Given the importance of telemedicine in the current public health situation and the possibility of integrating this strategy after the pandemic, it is essential to expand the knowledge about telemedicine use in different countries, types of technologies used and impacts on quality of care in the PHC. Thus, this study will map telemedicine use in PHC around the world during the COVID-19 pandemic and its impacts on quality of care.

**METHODS**

This is a scoping review protocol, a type of study designed to answer broad research questions with less restrictive selection criteria. The inclusion of scientific articles and the grey literature aim to map key concepts, types of evidence and research gaps, systematically synthesising the existing knowledge on a topic. The study will be developed according to Arksey and O’Malley and Levac et al., based on the Joanna Briggs Institute (JBI) manual, and guided by the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Extension for Scoping Reviews (PRISMA-ScR).

The following steps will guide this review: (1) formulation of research questions, (2) identification of relevant studies, (3) study selection, (4) data extraction and coding, (5) analysis and interpretation of results, (6) consultation with stakeholders.

**Step 1: formulation of research questions**

Three study questions were formulated and defined by consensus among authors using the PCC (Population–Concept–Context) mnemonic and outcomes of interest (O) (table 1).

Concepts and definitions anchoring the research questions are described in table 2.

**Step 2: identification of relevant studies**

**Search strategy**

A previous exploratory search was conducted on PubMed and Virtual Health Library (VHL) databases to identify the main Medical Subject Headings (MeSH) and Descriptors in Health Sciences (DeCS) related to the topic. The search strategy was developed based on this preliminary search by combining descriptors and keywords using the Boolean operators AND and OR, and it will be adjusted according to each database. A complete search strategy for PubMed database is included in online supplemental appendix 1, and an example of searching for grey literature on Google Scholar in online supplemental appendix 2. The detailed search strategy for all data sources (ie, white and grey literature) will be attached to the final scoping review.

**Table 1: Scoping review questions**

<table>
<thead>
<tr>
<th>Question</th>
<th>Population (P)</th>
<th>Concept (C)</th>
<th>Context (C)</th>
<th>Outcomes of interest (O)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(1) Which countries have used telemedicine in PHC due to the COVID-19 pandemic?</td>
<td>Countries in the world with PHC</td>
<td>Telemedicine</td>
<td>COVID-19 pandemic</td>
<td>Geographical mapping of telemedicine use</td>
</tr>
<tr>
<td>(2) What types of ICT in telemedicine have been used in PHC in the context of the COVID-19 pandemic?</td>
<td></td>
<td>ICTs</td>
<td>Classification of ICT types used</td>
<td></td>
</tr>
<tr>
<td>(3) What are the impacts of telemedicine on quality of care in PHC in the context of the COVID-19 pandemic?</td>
<td></td>
<td>Quality of care</td>
<td>Impact of telemedicine on patients' healthcare</td>
<td></td>
</tr>
</tbody>
</table>

ICT, information and communication technologies; PHC, primary healthcare.
Table 2  Key concepts for research questions

<table>
<thead>
<tr>
<th>Concept</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Teledicine</td>
<td>Provision of healthcare services by health professionals in situations in which distance is a critical factor using ICTs for research, continuing education of providers and health professionals, disease diagnosis, treatment, prevention and evaluation to promote individual and community health.14 15</td>
</tr>
<tr>
<td>Information and communication technologies</td>
<td>Integrated technological resources allowing access to information using telecommunication. It is the product of telecommunication, information technology and electronic media and serves as regulatory tools. It includes electronic devices used in healthcare, such as health information systems, electronic medical records, electronic prescriptions, mobile health, computers, electronic portals, emails, messages, cell phone applications, wearable devices (eg, smartwatches).16 36 37</td>
</tr>
<tr>
<td>Quality of care</td>
<td>It has three dimensions: technical (action choice accuracy and production), interpersonal (social and psychological relationships between healthcare providers and users) and organisational (conditions in which services are offered, including continuity of care, coverage, action coordination, access and accessibility).23–25 For this study: ▶ Technical dimension: Implementation of preventive and diagnostic measures and symptomatic treatment of mild COVID-19 cases in a timely manner (according to scientific protocols). ▶ Interpersonal dimension: Emotional support actions by healthcare providers to users and/or family members in situation of social isolation. ▶ Organisational dimension: Monitoring and follow-up actions of COVID-19 cases and continuity of care for patients without COVID-19.</td>
</tr>
</tbody>
</table>

Data sources

For a broader scope search, multidisciplinary health sciences databases and grey literature will be used. The following databases will be accessed: VHL, PubMed, Scopus, Web of Science, CINAHL and Embase. For the grey literature, searches will be conducted on Google Scholar (government guidelines, manuals, reports, documents, books and congress proceedings), preprint repositories (SciElo Preprints, bioRxiv and medRxiv) and specific databases for COVID-19 studies (WHO Global research on coronavirus disease, Cochrane Library, COVID-19 Open Research Dataset Challenge and Epistemónikos COVID-19).

Step 3: study selection

The study selection process for both the white and grey literature (identification, screening, eligibility and inclusion)32 will be presented in detail in the selection flow-chart of the review. These steps will be conducted by two independent reviewers (CRDVS and RHL), and, in case of disagreement, a third reviewer (SACU) will be consulted to reach a consensus.

Identified studies will be grouped in the Mendeley reference manager and duplicates will be removed. The Rayyan software will be used during title and abstract analysis to assist blinding of reviewers. Potentially relevant studies will be retrieved in full and exported to a Microsoft Excel (V.2016) database. Full texts will be analysed in detail according to eligibility criteria, and reasons for excluding studies will be recorded and reported in the review.

The selection process of studies from the grey literature will follow Godin et al.,33 and specific strategies will be conducted for searches on Google Scholar, preprint repositories and specific databases for COVID-19. The following terms will be combined: primary healthcare, teledicine, information and communication technologies, healthcare quality and COVID-19. The terms and number of studies retrieved will be recorded for each search strategy and the identified studies will follow the proposed selection steps mentioned above. Results from Google Scholar will be classified by relevance and the first 100 studies will be included in the screening process.33 Studies will be selected if data on at least one quality of care dimension (technical, interpersonal, organisational) are provided, as defined in table 2. Reference list of the identified studies will also be consulted for potentially relevant studies.

Before data collection, a pilot test will be conducted with all authors to reduce bias and ensure an aligned selection process: each author will select a random sample of 25 titles and abstracts, and screening will be performed according to eligibility criteria. Afterwards, the team discuss discrepancies and perform any necessary changes to criteria and definitions. Screening will initiate only after reaching an agreement of ≥75%,31 according to Fleiss’ Kappa statistics.31

Inclusion criteria

The following full-text studies focusing on teledicine in PHC during the COVID-19 pandemic, answering the study questions (table 1) and addressing at least one quality of care dimension (table 2) will be included: (1) primary studies, literature reviews, theoretical essays or brief communications; (2) grey literature, including preprints, guidelines, manuals, reports, government documents, books, dissertations, theses, and congress proceedings or other events of the academic community. Filters related to time will not be applied to searches since search strategies will already contain descriptors.
and terms related to the COVID-19 pandemic. Language filters will also not be used and an external translator will perform necessary translations. Searches will be updated 1 week before submitting the study to a scientific journal.

**Exclusion criteria**
This review presents a wide scope of research questions; therefore, only publications with inconsistent results or not answering the research questions will be excluded.

**Step 4: data extraction and coding**
Data related to the included studies will be extracted by two independent reviewers (CRDVS and RHL) using a data extraction form created based on the JBI template and adapted by the authors. The following information of interest will be retained: study description (title, first author, institution, year, objectives, study design, sample/participants, funding) and data answering the research questions (country that used telemedicine, ICT type, impact on quality of care) (online supplemental appendix 3).

A pilot test will be conducted with all authors and the extraction form filled with three studies to ensure that all necessary data will be retained properly. If necessary, the form will be refined to align the extraction process. Any changes will be reported in detail in the scoping review. Any divergences between reviewers during extraction process will be resolved by a third reviewer (SACU). When necessary, first authors will be contacted to request additional or missing data.

Google Earth V.7.15 and TerraView V.4.2.2 software will be used to identify and geocode the studies.

**Step 5: analysis and interpretation of results**
Data will be summarised quantitatively or qualitatively, as appropriate. For quantitative analysis, descriptive statistics (absolute frequencies and percentages) will be performed using SPSS software, V.24 (IBM). Qualitative analysis will be conducted using thematic analysis.35

This stage will be divided into data analysis, exposure of the results linked to research questions, and implications for other research and services. All results will be discussed with the relevant literature. Evidence synthesis will be presented using tables, diagrams, thematic maps, and, if possible, a meta-analysis will be conducted. A narrative summary reporting how results are related to the review purpose and research questions will accompany the mapped data.

**Step 6: consultation with stakeholders**
This will be the last step of the review.30 After analysing and interpreting, preliminary results will be presented to a group of experts in telemedicine and ICTs in PHC. The procedure will include invitation via email explaining the stakeholder participation and, if accepted, all materials will be sent to the panel of experts. Two web conferences will be scheduled to discuss results.

This strategy aims to share preliminary study findings (knowledge transfer and exchange), obtain potentially relevant studies not included in the initial search and develop effective dissemination strategies and directions for future studies.30

**Patient and public involvement**
In this protocol, there was no involvement of patients and the public. Stakeholder participation in the consultation stage is part of the scoping review dissemination plans. Patients will not be involved.

**Ethics and results dissemination**
In this study, secondary sources of information (ie, studies published in scientific journals and grey literature) will be included.

Human involvement will take place at the stakeholder consultation stage and will have the main purposes of defining dissemination strategies and sharing preliminary results of the study.

**Ethical approval**
Approved by the Research Ethics Committee of the Faculty of Health Sciences of Cariri, Federal University of Rio Grande do Norte (CAAE: 47473121.3.0000.5568).

Results will be disseminated through publication in an open access scientific journal, international congress in the field of collective health, and other media such as academic and community newspapers.

In case of changing this protocol after publication, information regarding changes, proper justification and dates will be provided.

The stages of the ongoing review are shown in **table 3**.

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**Table 3** Scoping review stages

<table>
<thead>
<tr>
<th>Stage</th>
<th>Start</th>
<th>Conclusion</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pilot searches to substantiate the review protocol and define the search strategy</td>
<td>2 September 2020</td>
<td>1 March 2021</td>
</tr>
<tr>
<td>Construction of the review protocol</td>
<td>11 September 2020</td>
<td>13 March 2021</td>
</tr>
<tr>
<td>Protocol registration in the Open Science Framework</td>
<td>18 March 2021</td>
<td>18 March 2021</td>
</tr>
<tr>
<td>Study selection</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Data extraction and coding</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Analysis and interpretation of results</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Consultation with stakeholders</td>
<td>No</td>
<td>No</td>
</tr>
</tbody>
</table>
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Contributors SaDCU proposed the study and coordinated the elaboration of the protocol. CRDVS and MFT developed the protocol. CRDVS, RHL, DiGBJ, MF-T, RAA and SaDCU participated in the discussion of the theoretical and methodological aspects of the study. CRDVS and RHL conducted the pilot searches to substantiate the search strategy. All authors reviewed the protocol and approved its final version for publication.

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Provenance and peer review Not commissioned; externally peer reviewed.

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REFERENCES