Telemedicine Infectious Diseases Consultations and Clinical Outcomes: A Systematic Review

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Background. Telemedicine use is increasing in many specialties, but its impact on clinical outcomes in infectious diseases has not been systematically reviewed. We reviewed the current evidence for clinical effectiveness of telemedicine infectious diseases consultations, including outcomes of mortality, hospital readmission, antimicrobial use, cost, length of stay, adherence, and patient satisfaction.

Methods. We queried Ovid MEDLINE 1946-, Embase.com 1947-, Scopus 1823-, Cochrane Database of Systematic Reviews (CDSR), Cochrane Central Register of Controlled Trials (CENTRAL), and ClinicalTrials.gov 1997- through August 5, 2019, for studies looking at clinical outcomes of infectious diseases in the setting of telemedicine use. We did not restrict by language or year of publication. Clinical outcomes searched included 30-day all-cause mortality, 30-day readmissions, patient compliance/adherence, patient satisfaction, cost or cost-effectiveness, length of hospital stay, antimicrobial use, and antimicrobial stewardship. Bias was assessed using standard methodologies. PROSPERO CRD42018105225.

Results. From a search pool of 1154 studies, only 18 involved telemedicine infectious diseases consultation and our selected clinical outcomes. The outcomes tracked were heterogeneous, precluding meta-analysis, and the majority of studies were of poor quality. Overall, clinical outcomes with telemedicine infectious diseases consultation seem comparable to in-person infectious diseases consultation.

Conclusions. Although in widespread use, the clinical effectiveness of telemedicine infectious diseases consultations has yet to be sufficiently studied. Further studies, or publication of previously collected and available data, are warranted to verify the cost-effectiveness of this widespread practice.

Keywords. clinical outcomes; infectious diseases consultation; mortality; systematic review; telemedicine.

Systematic review registration. PROSPERO CRD42018105225.

According to recent estimates, infectious diseases may be the third leading cause of death in the United States [1]. However, underserved and/or economically disadvantaged areas may not have access to infectious diseases (ID) physicians (up to 45% of US hospitals) to help treat these infections [2]. This is problematic because consultation with ID physicians significantly reduces mortality for numerous infections [3, 4]. Providing access to ID expertise in underserved/rural areas could substantially reduce mortality and improve clinical outcomes. As 51% of ID fellowship programs did not fill in 2015 [5, 6], access to ID expertise may be limited. With a shortage of ID physicians, it may not be possible for remote locations to employ an ID physician, that is, what is seen in current practice. Telemedicine could potentially expand ID expertise to underserved areas.

Telemedicine is widely used in many subspecialties. Studies show that telemedicine reduces mortality in progressive and intensive care units and in very low birth weight infants [7–9], but its effectiveness for important clinical outcomes in infectious diseases is lacking. To date, there has been no synthesis of evidence for the use of telemedicine for infectious diseases consultation. Our systematic review addresses this deficiency by answering the following question: In patients with infectious diseases, do telemedicine ID consultations improve the clinical outcomes of mortality, readmission, patient adherence/compliance, patient satisfaction, cost, cost-effectiveness, length of stay, antibiotic use, or antibiotic stewardship?

METHODS

Data Sources and Searches
A medical librarian (L.H.Y.) searched the literature for records including the concepts of infectious diseases, infection, antimicrobial stewardship, antibiotic stewardship, antifungal stewardship, antiviral stewardship, telemedicine and videoconferencing, and consultation/consult. The librarian created search
strategies using a combination of keywords and controlled vocabulary in Ovid Medline 1946-, Embase 1947-, Scopus 1823-, Cochrane Database of Systematic Reviews (CDSR), Cochrane Central Register of Controlled Trials (CENTRAL), and Clinicaltrials.gov 1997-. All search strategies were completed in November 2018, then updated in August 2019. The protocol was registered with the international prospective register of systematic reviews (PROSPERO), in accordance with PRISMA-P guidelines (PROSPERO CRD42018105225) [10]. The protocol for this systematic review has been previously described [11].

Aim
Our goal was to assess the effectiveness of telemedicine ID consultation for a range of clinical outcomes (enumerated below) as compared with either (1) no ID consultation or (2) other modalities of ID consultation (eg, in person). Clinical outcomes considered included 30-day all-cause mortality, readmission within 30 days of discharge from an initial hospitalization with an infection, patient compliance/adherence, patient satisfaction, cost or cost-effectiveness, length of hospital stay, antimicrobial use, and/or antimicrobial stewardship.

There were no language or year of publication restrictions. Translation of non-English-language abstracts was undertaken, as required, though ultimately all full-text articles were in English. Conference abstracts were excluded if sufficient outcome and bias data could not be extracted. REDCap was used for data entry.

Study Selection
After removal of duplicate results, titles and abstracts were reviewed for relevance to the research question by J.P.B. See Figure 1 for the study flowchart from title/abstract review to final inclusion. Articles were excluded if any of the following conditions was met: (1) infections not studied, (2) no consultations, (3) consultations not performed for infection, (4) no telemedicine, (5) noninterventional (eg, viewpoint articles, commentaries, etc.), (6) infectious diseases outcomes indistinguishable from other consultations, (7) no prespecified outcomes of interest tracked, or (8) abstract only with insufficient methodological or results reporting. Studies that were not excluded underwent independent, blinded, full-text review by J.P.B. and G.A.C.

Definitions
Telemedicine was defined as remote clinical services administered using a technological medium. This included face-to-face video chat (physician-to-physician or physician-to-patient), voice chat after review of electronic health records, or electronic health record documentation after remote chart review without direct voice or video contact with physician or patient. Antibiotic stewardship was quantified as either antibiotic costs or antibiotic appropriateness, as judged by the authors of the individual studies.

Data Extraction
In a blinded fashion, 2 authors (J.P.B. and G.A.C.) independently extracted data from full-text articles. Data extracted included study quality, clinical or system-level outcome tracked, percent change or proportion experiencing each clinical outcome, numbers of patients, age group, consultant specialty, type of telemedicine, study location, whether infection was confirmed by laboratory results, and type and risk of bias.

Data extraction disputes were settled by a third reviewer (S.A.F.—also blinded), and in cases requiring further adjudication, a group session of all 3 reviewers was convened.

Quality Assessment
Risk of bias was independently reviewed by 2 reviewers (J.P.B. and G.A.C.) in a blinded fashion. Disputes were resolved by a third reviewer (S.A.F.—also blinded), and in cases of continued disagreement, the 3 reviewers met for adjudication. Bias determination was guided by the Cochrane Consumers and Communication Review Group Study Quality Guide or Newcastle-Ottawa scale [12, 13]. Using the Newcastle-Ottawa Scale, the case–control and cohort studies were given star ratings.
in 3 categories—Selection (maximum 4 stars), Comparability (maximum 2 stars), and Outcome (maximum 3 stars)—with a maximum score of 9 stars [13]. The quality of case–control and cohort studies was adjudicated based on previously published guidance [14]: good quality: Selection ≥3 stars AND Comparability ≥1 stars AND Outcome ≥2 stars; fair quality: Selection 2 stars AND Comparability ≥1 stars AND Outcome ≥2 stars; poor quality: Selection ≤1 Star OR Comparability 0 stars OR ≤1 stars.

Analysis

Per protocol [11], prespecified subgroup analyses included outcomes by age (children <18 years vs adult ≥18 years), telemedicine consultant being ID trained or not, infection type, type of telehealth/telemedicine intervention (eg, face-to-face, asynchronous, etc.), study location (US vs non-US), number of ID consultations (ie, days physician interacted with patient/provider), and culture- or laboratory-confirmed infection vs presumed infection. Due to the limited number of studies, qualitative/narrative synthesis was performed.

RESULTS

A total of 1328 results were found using our initial search strategy, which was completed in November 2018. A total of 284 duplicate records were identified using Endnote’s automatic duplication finder, and another 31 duplicates were removed by manual review, leaving 1013 unique citations in the project library. One additional study was identified after reviewing references of full-text article reviews, for a total of 1014 search results. We updated our search on August 5, 2019. With this update, an additional 140 results were found, leaving a total of 1154 results to be reviewed. Fully reportable searches can be found in Appendix 1.

Of the 1154, none of the clinical trials identified only from ClinicalTrials.gov (n = 21) had available results. Of the remaining excluded articles, the reasons for exclusion are listed in Figure 1 and Supplementary Table 1. A total of 18 articles were relevant to the research question and underwent full-text review. From these articles, clinical outcomes tracked included 30-day mortality after an infection (16.7%, n = 3), readmission within 30 days after discharge from the initial hospitalization with an infection (5.6%, n = 1), patient compliance/adherence (11.1%, n = 2), patient satisfaction (50.0%, n = 9), cost or cost-effectiveness (22.2%, n = 4), length of stay (27.8%, n = 5), and antimicrobial use (27.8%, n = 5). Meta-analysis was not performed due to the low number of studies with any 1 outcome.

Biases/Quality Assessment

There were 2 (11.1%) randomized clinical trials, 2 case–control studies (11.1%), and 14 (77.8%) cohort studies. Both randomized controlled trials had high risk of bias in 3 categories and unclear risk in 2 others [15, 16]. Using the Newcastle-Ottawa Scale, both the case–control studies and 8 cohort studies were rated as poor quality. The remaining 6 cohort studies were rated as good quality (Supplementary Table 2).

Study Characteristics

Most studies were performed only in adults (n = 13, 72.2%). Consultant specialty was infectious diseases in only 38.9% (n = 7). Infections studied included pneumonia (n = 4), urinary tract infection (n = 5), sepsis (n = 3), bacteremia (n = 3), endocarditis (n = 2), skin and soft tissue infections (n = 3), upper respiratory infections (n = 4), and other (n = 12). More than 1 infection type could be studied in each article.

The most common type of telemedicine was face-to-face videoconferencing with the patient in 72.2% (n = 13), followed by telephone only (16.7%, n = 3), physician-to-physician only (5.6%, n = 1), and 1 study in which the telemedicine type could not definitively be determined.

Just over half (n = 10) were based in the United States, and 8 in other countries (Europe = 4, Australia = 2, Asia = 1, Canada = 1). Infections were confirmed by culture in 61.1% (n = 11) of studies.

Clinical Outcomes

Patient satisfaction with telemedicine was the most commonly reported outcome, and the percentage of patients satisfied with telemedicine was above 97% in 6/7 studies [17–22], with 1 study reporting patient satisfaction of 69% (Table 1) [16]. Two additional studies reported patient satisfaction but provided a mean satisfaction score without a numerator and denominator for the number of patients reporting the outcome, though the mean score in both studies was indicative of high satisfaction [23, 24].

Mortality was higher in the telemedicine group in 2 studies and lower in the other 2 studies reporting this outcome (range for all studies, 0%–22%) [15, 25–27], with 1 study reporting 90-day instead of 30-day mortality (higher mortality in the control group) (Table 1) [27]. Only 1 of these studies was statistically significant, with lower mortality in patients receiving in-person rather than telephone-only ID consults [26]. Length of stay was shorter in the telemedicine group in 4/5 studies [24–27] and equivalent in 1 study (range, 2.6–30 days) (Table 1) [28].

Readmission and adherence/compliance were similar between telemedicine and nontelemedicine groups (Table 1) [15, 19, 27]. Costs were lower in the telemedicine groups, but based on projections that may not be generalizable [15, 24, 28, 29]. Antibiotic use was similar between telemedicine groups and controls [25, 27, 30–32].

DISCUSSION

Based on the available, albeit limited, evidence, telemedicine ID consultation seems comparable to standard of care for the
clinical outcomes of mortality, length of stay, readmission, adherence, cost, and antimicrobial use. However, there were few relevant studies tracking our prespecified clinical outcomes, and the majority were of poor quality. Without more robust data quality and availability, it is difficult to draw any firm conclusions.

Few studies have been published demonstrating effectiveness of telemedicine for infectious diseases consultation. Infection types that have been studied, as well as the settings in which telemedicine has been used, have been varied. As telemedicine continues to expand, clinicians and researchers should consider publication of their already existing data to document the clinical effectiveness needed to validate this model of care and costs. In addition, researchers must report their processes of telemedicine ID implementation so that what works in 1 study can be applied more broadly, understanding that adaptations will likely be required. In this rapidly blossoming field, we must publish best practices using standardized reporting for effective and implementable telemedicine ID consults so that our patients reap maximum benefits.

Our review is limited in scope by our chosen clinical outcomes. Among the excluded studies (data not shown) were studies looking at the use of telemedicine to care for patients with hepatitis C or HIV (among others). These studies are important and relevant to the telemedicine ID consultation landscape, but did not track our prespecified clinical outcomes. The clinical outcomes we chose are of interest to inpatient settings (mortality, readmission, length of stay) and administrators (cost, patient satisfaction, readmission, antimicrobial use, mortality). A summary of the currently available data may help in the adoption of inpatient telemedicine ID services by inpatient physicians and administrators, should the data become more robust.

Infectious diseases consultation can save lives, and with almost half of US hospitals without ID physician access [2], telemedicine has great potential to fill this gap. Before its wide adoption, it should be robustly evidence-based. Important questions in this area are related to which type of telemedicine is required, how frequently telemedicine visits must be performed (eg, daily, 1-time), and what level of infectious
diseases physician involvement is optimal to achieve the best clinical outcomes. For example, a recent study showed that an algorithm-based care model for *Staphylococcus aureus* bacteremia achieved noninferior outcomes to usual care [33]. With this in mind, one must ask what the minimum unit of efficacy is for infection management. Whether that is in-person consultation, telemedicine consultation, Extension for Community Healthcare Outcomes (ECHO)–like models, algorithm-based care, or another care model has yet to be determined. In addition, it is unknown whether certain infections will require different levels of ID input for optimal outcomes. Notably, curbside consultations are often inaccurate and potentially harmful [34], which may have implications for care delivery methods (eg, telephone consultation only). Further studies are required.

Many questions remain to be answered for telemedicine ID consultation, including reimbursement, as state-to-state differences in telemedicine coverage remain a barrier to implementation. As the field of telemedicine continues to grow, these questions must be addressed to provide the best and most efficient care for patients. Societal and technological barriers such as access to high-speed Internet and video quality have limited telemedicine’s usefulness in the past. Although Internet access issues persist in some rural areas, progress has been made and video quality has improved. High-priority areas of telemedicine research include publishing data that are already being collected as part of routine clinical care, understanding the use of telemedicine in rural/underserved settings and how it reduces barriers to care and reduces the health care disparities therein, and determining how to most efficiently deliver care (eg, face-to-face vs e-consult, etc.). Telemedicine ID consultation may be a way to reduce inequities and treatment disparities for rural/economically disadvantaged patients, and from the perspective of primum non nocere, we must be sure that what we are doing is what is best.

**Supplementary Data**

Supplementary materials are available at Open Forum Infectious Diseases online. Consisting of data provided by the authors to benefit the reader, the posted materials are not copyedited and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

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