

Telemedicine in rheumatology: A mixed methods study exploring acceptability, preferences and experiences among patients and clinicians.

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Abstract

Objectives

The Covid-19 pandemic necessitated a rapid global transition towards telemedicine; yet much

remains unknown about telemedicine's acceptability and safety in rheumatology. To help address

this gap and inform practice, this study investigated rheumatology patient and clinician experiences

and views of telemedicine.

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Methods

Sequential mixed methodology combined analysis of surveys and in-depth interviews. Between and within-group differences in views of telemedicine were examined for patients and clinicians using t-tests.

Results

Surveys (Patients n=1,340, Clinicians n=111) and interviews (Patients n= 31, Clinicians n=29) were completed between April 2021 and July 2021. The majority of patients were from the UK (96%) and had inflammatory arthritis (32%) or lupus (32%). Patients and clinicians rated telemedicine as worse than face-to-face consultations in almost all categories, although >60% found it more convenient. Building trusting medical relationships and assessment accuracy were great concerns (93% of clinicians and 86% of patients rated telemedicine as worse than face-to-face for assessment accuracy). Telemedicine was perceived to have increased misdiagnoses, inequalities and barriers to accessing care. Participants reported highly disparate telemedicine delivery and responsiveness from primary and secondary care. Although rheumatology clinicians highlighted the importance of a quick response to flaring patients, only 55% of patients were confident that their rheumatology department would respond within 48 hours.

Conclusion

Findings indicate a preference for face-to-face consultations. Some negative experiences may be due to the pandemic rather than telemedicine specifically, although the risk of greater diagnostic inaccuracies using telemedicine is unlikely to be fully resolved. Training, choice, careful patient selection, and further consultation with clinicians and patients is required to increase telemedicine's acceptability and safety.

Key messages:

- 1. Concerns about telemedicine included: reduced assessment accuracy; impaired medicalrelationship building; and increased inequalities for vulnerable and/or disadvantaged patients.
- 2. Barriers to patient-initiated contact, including 'gate-keeping' administrators and nonresponse, are limiting emergency access.
- 3. Telemedicine is convenient and acceptable for some rheumatology patients. Careful selection and choice are key.

Key words: Telemedicine, rheumatology, patient-physician interactions, digital technology in medicine, pandemic, mixed-methods, rare autoimmune rheumatic diseases

Introduction

The Covid-19 pandemic necessitated a rapid reorganisation of patient care to predominantly telemedicine in the UK¹ and elsewhere^{2,3,4}. This trend seems set to continue⁵, with support from governments and health-services⁶. NHS England's 2021/2022 planning guidance, for example, states that at least 25% of outpatient appointments should be by telephone or video⁶. However, telemedicine's suitability, acceptability and safety in rheumatology has yet to be ascertained. The majority of rheumatology studies have focused on rheumatoid arthritis, and there is limited evidence relating to the rare autoimmune rheumatic diseases^{7,8} such as lupus or systemic sclerosis, which can have life-threatening⁹, yet less visible¹⁰ manifestations.

McDougal et al's tele-rheumatology systematic review found that most studies reviewed 'demonstrated a high risk of bias'¹¹; and studies have produced conflicting results. Some report high levels of patient satisfaction^{12,13} and suggest that telemedicine could be as effective as face-to-face for autoimmune rheumatic diseases¹⁴. However, there are concerns regarding the use of

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telemedicine for diagnosis and assessment of disease activity¹⁵, particularly as a result of lack of physical examinations¹⁶. Studies to date have tended to be limited by sample size¹⁴, only reporting the views of one side of the medical relationship and/or only including patients from the investigators' hospital^{13,17}, which could increase the risk of bias.

The pandemic-initiated extensive use of telemedicine has highlighted the need for detailed research ahead of more permanent digital service delivery. Our study therefore elicited the views and experiences of clinicians, and patients with a wide range of inflammatory rheumatic diseases.

Methods

Participants and design

A pre-tested survey for rheumatology patients was made available online in April 2021 using Qualtrics, via multiple disease support groups. A similar online survey for rheumatology clinicians was disseminated through rheumatology networks, including the British Society for Rheumatology, in May/June 2021. This telemedicine sub-study is part of a pre-registered longitudinal multi-stage trial (ISRCTN-14966097, including a statistical analysis plan with Covid-related additions registered March 2021). Ethical approval was obtained through the Cambridge Psychology Research Committee. Respondents gave informed written (electronic) consent on the survey and/or verbally recorded consent prior to interviews.

Inclusion criteria: Patients: age ≥18 years, reporting a diagnosis of an autoimmune inflammatory rheumatological condition on their clinic letters and having received at least one telemedicine appointment. Clinicians: UK-based clinicians involved in the care of rheumatology patients (including consultants, registrars, specialist nurses and General Practitioners (GPs)).

Questions elicited demographic, and quantitative (predominately using Likert scales comparing telemedicine with face-to-face (F2F)) and qualitative responses about telemedicine views and

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experiences. Medical security was measured on a scale of 0-100 with 0 signifying 'completely medically abandoned' to 100 signifying 'completely medical secure'. Interviewees were largely purposively selected from the questionnaires to ensure a range of characteristics and views. Interviews were semi-structured, continued until thematic saturation was reached and were conducted by M.S, an experienced, qualitatively-trained researcher. Interviews lasted for 0.5-3 hours and were transcribed verbatim.

Analysis

This study used multi-stage sequential mixed methods. Interview data was collected both before survey design to ascertain patient and clinician priorities for research directions, and also after initial analysis (using SPSS v.26) of quantitative data to further explore and explain quantitative findings. Both types of data were given equal priority and the use of mixed methods reduced the inherent weaknesses in each individual method. Mean differences between patients and clinicians, and within-groups differences, were tested for statistical significance using t-tests. Pearson's or Spearman's rank were used for correlations as appropriate. Thematic analysis¹⁸ was used for the interviews and the qualitative data from the surveys, using NVivo12 to assist in managing and coding the qualitative data. The coding frame was developed by MS, RH and MB, with coding agreed by FN, EL, PH and CG, to improve reliability. Emerging themes were discussed and agreed by the wider team, including patient representatives and clinicians. Validity was strengthened by triangulation of quantitative and qualitative results¹⁹, which were combined at all stages from planning through analysis to reporting, consideration of cases that deviated from the emerging conclusions²⁰ and member checking²¹. The criteria for reporting qualitative research (COREQ)²², detailed methodology and limitations are included in Supplementary Data S1, available at Rheumatology online. Further statistical analysis is available in Supplementary Data S2, available at *Rheumatology* online.

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Ethical Approval: Ethical approval was obtained through the Cambridge Psychology Research Committee: PRE.2019.099: approval for original trial; PRE.2020.089: approval for COVID-19 related additions and amendments.

Trial registration: This telemedicine study is part of a pre-registered longitudinal multi-stage trial, the LISTEN study (ISRCTN-14966097), with later Covid-related additions registered in March 2021, including a pre-registered statistical analysis plan.

Results

A total of 1,340 patients and 111 clinicians completed the surveys; 60 (31 patients and 29 clinicians) were interviewed (table 1). The four most commonly reported rheumatological diseases were inflammatory arthritis (32%), lupus (32%), Sjogren's (9%), and systemic sclerosis (8%). The survey was open to clinicians working in the UK and patients of any nationality, but patient respondents were largely from the UK (96%). Numerical values given are from survey quantitative data. Within text quotes were not given participant numbers if the term used was given by multiple participants.

Table 1. Participant characteristics

Characteristic	Patient survey	Patient	Clinician survey	Clinician
	(N=1340)	interviews (N=31)	(N=111)	interviews (N=29)
Age				
<30	222 (17%)	4 (13%)	5 (5%)	0
30-39	247 (18%)	3 (10%)	19 (17%)	9 (31%)
40-49	406 (30%)	8 (26%)	39 (35%)	5 (17%)
50-59	308 (23%)	9 (29%)	37 (33%)	11 (38%)
60+	155 (12%)	7 (23%)	11 (10%)	4 (14%)
Gender				
Female	1275 (95%)	27 (87%)	77 (69%)	11 (38%)
Male	59 (4%)	4 (13%)	34 (31%)	18 (62%)
Other/undisclosed	6 (<1%)	0	0	0
Country/region				
England	1060 (79%)	17 (55%)	87 (78%)	24 (83%)
Scotland	112 (8%)	4 (13%)	13 (11%)	2 (7%)
Wales	68 (5%)	8 (26%)	9 (8%)	1 (3%)
N. Ireland	36 (3%)	0	2 (2%)	2 (7%)
US/Canada	30 (2%)	1 (3%)	0	0

Europe	16 (1%)	1 (3%)	0	0
Other	18 (1%)	0	0	0
Disease				
Inflammatory (predominantly Rheumatoid) arthritis	424 (32%)	7 (23%)		
Lupus	419 (32%)	12 (39%)		
Sjögrens	119 (9%)	2 (6%)		
Systemic sclerosis	100 (8%)	3 (10%)		
Vasculitis	47 (4%)	1 (3%)		
UCTD	45 (3%)	3 (10%)		
PMR	44 (3%)	2 (6%)		
MCTD or two or more inflammatory rheumatic diseases	91 (7%)	1 (3%)		
Other inflammatory rheumatic disease	47 (4%)	0		
Time since diagnosis				
< 1 year	80 (6%)	3 (10%)		
1-2 years	166 (12%)	5 (16%)		
3-5 years	268 (20%)	5 (16%)		
6-9 years	253 (19%)	9 (29%)		
10+ years	568 (42%)	9 (29%)		
Unsure or missing	5 (<1%)	0		
Clinician Role				
Rheumatology consultant			72 (65%)	13 (45%)
Rheumatology registrar			13 (12%)	5 (17%)
Rheumatology nurse			19 (17%)	5 (17%)
GP			0 (not surveyed)	5 (17%)
Other speciality			7(6%)	1(7%)

Five main telemedicine themes were identified: 1) Effect on the patient-clinician relationship; 2) Greater risk of inaccurate assessments; 3) Convenience, technology and logistical considerations; 4) Barriers to safe and effective telemedicine; and 5) Cost-cutting, clinical need or choice?

Theme 1: Effect on the patient-clinician relationship

Significantly more (p=0.02) clinicians (90%) than patients (69%) felt that telemedicine was worse than face-to-face for building a trusting relationship (Figure 1a). The majority of clinicians discussed how non-verbal communication, including *'reassuring'* touch, was important in building trust and/or rapport. Many patients used terminology - such as feeling like *'a statistic'* or *'uncared for'* - indicative of finding telemedicine more impersonal than in-person consultations. However, a substantial

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number of participants reported that positive relationships continued with telemedicine; and wellestablished medical relationships were felt to be protective against the limitations of remote appointments.

Around 50% of clinicians and patients rated clinicians listening as worse with telemedicine, and <10% rated it better than with face-to-face. Many patients felt telemedicine followed an impersonal *'tick list'* and felt more *'rushed'*. Patients' ratings for telemedicine were significantly lower (mean difference, 0.49, p<0.001) than clinicians for time available to discuss patient concerns (Fig 3a), and many reported telephone consultations lasting <5 minutes. Other patients expressed disappointment regarding changes in their clinician's behaviour using telemedicine:

The doctor, who is normally quite sympathetic, seemed very cold...very perfunctory and I felt he sounded rather irritated. I didn't feel very cared for at the end of the call, despite getting the outcome I felt I needed (patient 56, RA)

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Some clinicians surmised that pandemic-induced backlogs might be temporarily reducing time available and patience during discussions, whilst others felt that telemedicine itself hindered displaying empathy or their ability to focus on the patient:

When I am in clinic I am 100% focused on the patient I have in front of me...but on the telephone I really struggle... we have less time. I am always watching the watch, I feel much more tired than with face to face (Clinician 23, Consultant)

Theme 2: Greater risk of inaccurate assessments

Patients (86%) and clinicians (93%) felt that telemedicine was worse than face-to-face in terms of accuracy of assessment (Fig 1b). Misdiagnoses and other inaccuracies were frequently reported and often attributed to the absence of examinations and visual cues:

My rheumatologist cannot see/hear how I move, look at my skin, eyes, hair, hands, bones, how I am...I was diagnosed with something over the phone, which I know isn't right, and it's getting worse (Patient 293, UCTD)

Compounding these problems was the cancelling of tests that could aid diagnoses. Patients reported having at least one blood (40% of patients) and/or other (48%) test (e.g. imaging) cancelled between March 2020 and March 2021. During that time, routine medication monitoring was restricted to protect patients from Covid-19 infection, but it remains limited, often due to continuing disagreements between primary and secondary care as to responsibility for testing when patients are being reviewed remotely:

Secondary care has gone to much more telephone and requesting that GPs do the blood tests...I'm not sure what that abnormal result means, it feels like we're doing secondary care's job for them. It is less safe for the patient (Clinician 29, GP)

The risk of diagnostic errors from lack of examinations and tests was further increased by frequent under-playing of symptom severity by rheumatology patients. Willingness to report symptoms, including mental health (MH) symptoms, was reported as worse/much worse with telemedicine by approximately 45% of patients and clinicians. Clinicians also found it more difficult to identify MH concerns remotely:

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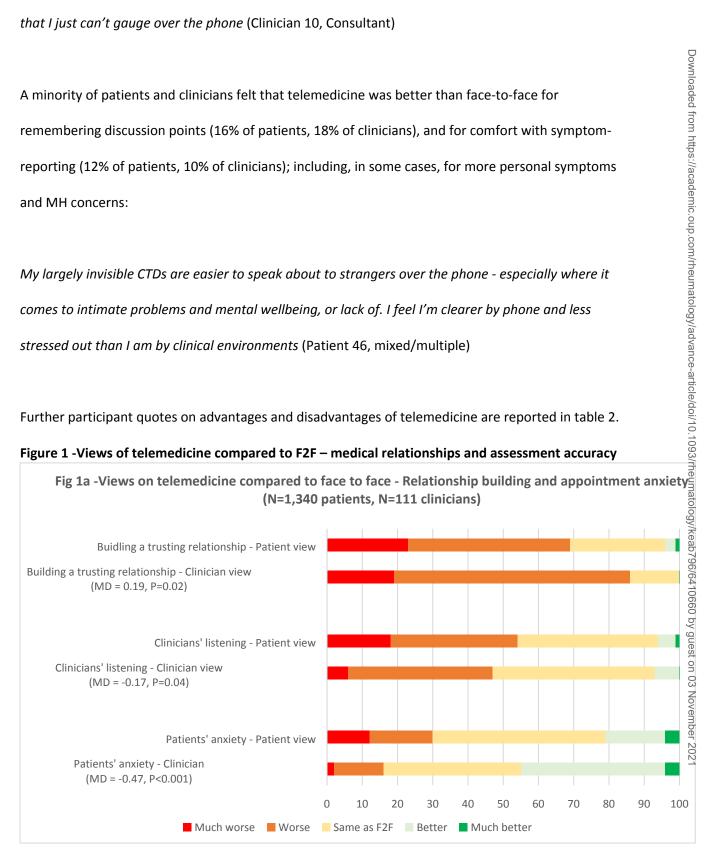
I can tell from somebody's affect and demeanour, and how they look and how they're different from how they've been previously and how they respond, you know those biological features of depression that I just can't gauge over the phone (Clinician 10, Consultant)

A minority of patients and clinicians felt that telemedicine was better than face-to-face for remembering discussion points (16% of patients, 18% of clinicians), and for comfort with symptomreporting (12% of patients, 10% of clinicians); including, in some cases, for more personal symptoms and MH concerns:

My largely invisible CTDs are easier to speak about to strangers over the phone - especially where it comes to intimate problems and mental wellbeing, or lack of. I feel I'm clearer by phone and less stressed out than I am by clinical environments (Patient 46, mixed/multiple)

Further participant quotes on advantages and disadvantages of telemedicine are reported in table 2.

Figure 1 -Views of telemedicine compared to F2F – medical relationships and assessment accuracy



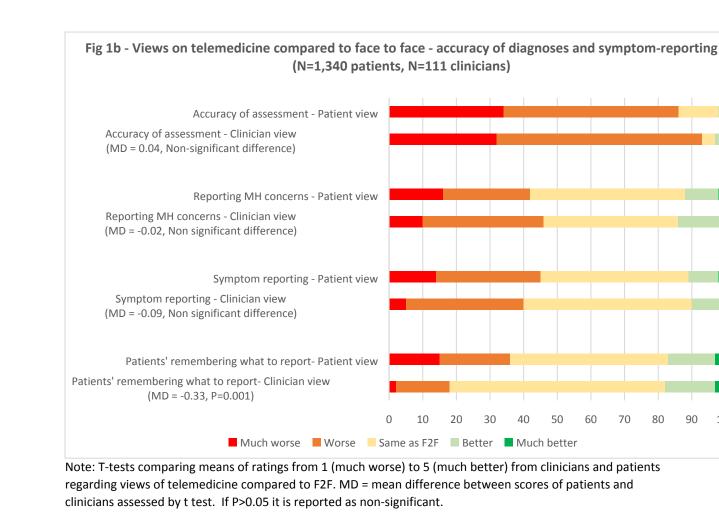


Table 2. Illustrative participant quotes on the advantages and disadvantages of telemedicine

Category	Advantages of telemedicine	Disadvantages of telemedicine
Building a trusting relationship	It's nice that my GP has called me. This way I didn't have to go to her Practice. She has also called me to check if I'm having flares. Small things like this really kept me going during lockdown (Patient 1063, lupus)	It's definitely harder to build trust over the phone, and you're picking up that they're not saying everything, so much of that for me on my side is that I use very open body language, eye contact, that kind of stuff, and showing empathy (Clinician 11, Registrar)
Clinicians' listening and taking symptoms seriously	Medics listened to symptoms because they couldn't see me. More so than in person. (Patient 517, Sjögrens)	All the time he was interrupting and my words were falling on deaf ears 'How's your skin?' Me, 'starting to flare'. Again no advice or reaction I felt like I was on a very long patient list he had to get through and he was taking no prisoners. I decided I didn't want to see him again and was greatly upset by his abruptness and attitude (Patient 23, lupus)
Patients' feeling anxious due to previous adverse	I get huge anxiety just walking into the doctor's office because I feel like I'm not going to be heard. Also, I've had so many bad experiences, I don't feel like doctors are interested enough. So, in that regard,	It is always different on the phone whereas if you see them you can see how they're actually feelingoften feel like I'm not being believedmakes me report it less on the phone definitely. (Patient 80, lupus)

medical experiences	<i>telemedicine is awesome. I'm in my home, on my territory and they can't harm me</i> (Patient 892, RA)	
Accuracy of assessment	No advantages in this category specified by clinicians or patients	Telemedicine sucks and kills people. Might be good for administrative stuff, like questions about medications etc but dangerous for if you are actually unwell. Easy to dismiss a patient ov the phone and impossible to physically exam them. And idiotic to make the patient and their family behave like medical professionals (Patien 836, mixed/multiple)
Reporting of symptoms, including mental health	I like themhonestly I feel more listened to and less dismissed Maybe it's changed how I present myself too as I'm more confident in my own space and not so aware of being judged visually (Patient 890, Systemic sclerosis)	It's quite easy to hide how you're really feeling. I've done this when my anxiety was having a wobbleneeded more support than I let on. It's much easier to put a brave face on when on a phone call (Patient 578, dermatomyositis)
Remembering what to report	It's helpful being able to make notes before and during the appointment when it's by phone so you don't forget to ask about all pertinent issues I don't feel confident doing that in a face-to-face appointment. I think doctors hate it when you produce a list at an appointment (Patient 310, RA)	It's harder for them to remember they're thinking about what's going on in the background and in the house. They don't actual then think so much and focus than if they were sitting in your office and there were no distractionsAnd they are not then as mentally prepared (Clinician 5, GP)
Convenience	We are a tertiary centre and most of our patients travel for hours to get to us and it does seem bonkers at times when they do that and stay for 5 minutesclearly a role for patients like that and I know a lot of them have really liked it. (Clinician 11, Registrar)	<i>If we knew a time slot it would really improve telemedicine. I'm a teacher and can't take a call in class</i> (Patient 1385, vasculitis)

Theme 3: Convenience, technology and logistical considerations

Over 60% of patients and clinicians considered telemedicine more convenient than face-to-face (Figure 2), often citing safety from infection, no travelling, and reduced waiting times, as the main

advantages (Table 3a):

I am very glad that telemedicine has become an option as it not only makes me feel more relaxed

and safer, but I often have great difficulty getting to hospital (Patient 635, lupus)

Clinicians significantly over-estimated the convenience for patients compared with patient views

(mean clinician rating of 3.86, mean patient rating of 3.49, mean difference=0.37, p=0.001), and

many patients reported that they weren't given appointment times for telemedicine in the same

way as for face-to-face, leading to greater inconvenience for some. In addition, remote

appointments did not always save clinicians' time (Fig. 2b). Telemedicine involves more

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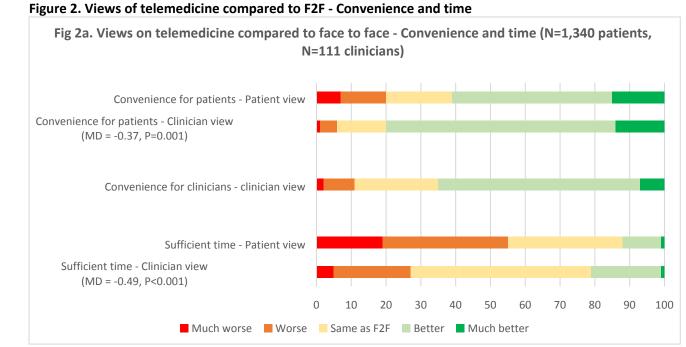
administration, including sending blood forms and prescriptions that would have been handed to the patient in a face-to-face appointment. Patients were often '*double-seen*' by clinicians who arranged a face-to-face appointment for any they were concerned about. Administrative errors meant that patients were sometimes not informed as to whether their appointment was face-to-face or telemedicine. Although some clinicians reported an improved work-life balance through remoteworking, others found their workload expanding as a result of the increase in direct patient communication (such as by email).

NHS technology and organisation were sometimes spoken of in strongly negative terms, such as being 'embarrassingly primitive' (Clinician 82, Consultant); although several clinicians reported rapid pandemic-induced improvements in technology. Despite sometimes 'fierce opposition' (Clinician 4, Consultant) to change initially from some clinicians, hospitals with existing integrated technology for communication between clinicians and patients had an easier transition to telemedicine and greater patient satisfaction:

The addition of the "MyChart" app...has really helped me. I can send a message without battling to pick up the phone, I can attach photos which can then be shown to my rheumatologist, and I receive all my blood test results quickly too (Patient 825, mixed/multiple)

Clinicians and patients felt that the medical relationship and the quality of care would benefit from improved telemedicine skills for patients: *once my daughter had taught me how to email photos I sent some flare pictures in* (Patient 1424, Sjogren's), and also for clinicians:

I have taught telemedicine myself as a clinical subject. What is apparent in the current situation is the lack of preparedness in the NHS to train clinical staff in telemedicine in terms of: demonstrating empathy, active listening, summarising and checking, formulating a shared management plan and provision of signposting, care and worsening advice (Patient 1250, lupus)



Note: T-tests comparing means of ratings from 1 (much worse) to 5 (much better) from clinicians and patients regarding views of telemedicine compared to F2F. MD = mean difference between scores of patients and clinicians assessed by t test. If P>0.05 it is reported as non-significant

Fig 2b. Common clinician views of advantages and disadvantages of te	lemedicine for convenience
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Category	Advantages of telemedicine	Disadvantages of telemedicine
Work/life balance	Able to work from home intermittently has lessened the stress of daily travel (Clinician 81, Nurse)	Inundated with emails from patientsyesterday spent 2 or 3 hours just dealing with patient emails. (Clinician 4, Consultant)
Time	More efficient use of time for self and patient when they are in remission (Clinician 53, Nurse)	Sometimes we double see patients because we're trying to deal with the problem on the telephone and then you end up trying to schedule them again to come in, and you're talking about pressured waiting lists already (Clinician 1, Registrar)
Productivity	Makes things easier for clinicians as we can more easily multi-task, check resultswhile speaking to the patient without having to make eye contact (Clinician 77, Registrar)	If you don't answer then they keep callingkeep on being interrupted by all these phonecalls then answering an email and then lots of calls that nobody is takinginterrupts me when maybe I'm maybe checking bloods and things can then go a bit wrongit's distractions all the time (Clinician 23, Consultant)

Theme 4: Barriers to safe and effective telemedicine

Although most clinicians acknowledged that problems had occurred with patient-initiated access

during earlier stages of the pandemic, the majority felt that their own department/GP surgery was

currently responding promptly. Several highlighted the importance of a quick response to patients experiencing flares:

If a patient phones up a department, even if we're working remotely, a lupus patient should be responded to on the same day, there's no doubt about that (Clinician 25, Consultant)

A minority of patients reported that increased use of telemedicine had ensured quicker responses:

The GPs are much more efficient using telephone appointments and when I've needed them they've seemed much more responsive and engaged. The NHS has benefitted from changing how they work (Patient 890, Systemic sclerosis)

However, emergency access remained highly variable and many patients felt *'insecure'* due to continued difficulties in obtaining emergency care/advice. Overall patient medical security ratings (from 0 = totally medically abandoned to 100 = totally medically secure) were lower for patient recalled pre-pandemic levels (66.7) compared to during (48.1) the pandemic (mean difference 18.6, p<0.001). Differences between disease groups' ratings of medical security increased during the pandemic. For example, recalled pre-pandemic medical security ratings were 68.9 for participants with inflammatory arthritis and 66.0 for participants with lupus (with a non-significant mean difference of 2.9, p=0.06). These had dropped to 51.8 (inflammatory arthritis) and 46.3 (lupus) in April 2021 (MD of 5.5, p=0.004). Medical security was lowest in Sjogren's patients (42.2 in April 2021).

Great difficulties in accessing support were frequently reported, sometimes from the departments that clinicians felt were responding promptly:

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Waiting for a call back after 4 voicemails...feel sad and scared knowing that when I really need medical help I have no-one (Patient 132, lupus)

Administrative staff triaging and 'gate-keeping' was particularly disliked; and there were reports of being made to feel 'like I was making a fuss' or being refused access, sometimes with life-threatening consequences:

You have to go through reception, she...didn't think I should bother GP... hung up on me...it was 3 or 4 days that she agreed that actually a doctor did need to see me...my lovely GP who was like it's too late, straight to A&E...kidney infection. I was in hospital for a week... If I had seen the GP in real life this would not have happened (Patient 9, lupus)

Perceptions of continued barriers to prompt care were also reflected in the quantitative findings, with for example only 48% being confident that a GP would respond within 24 hours, and only 55% confident that their rheumatology department would respond within 48 hours. Patients with inflammatory arthritis were significantly more confident (p=0.002) of getting a prompt response from their rheumatology department than those with lupus. Restricted methods of access were felt to disproportionately impact the most disadvantaged:

We've had some local Practices only allowing contact through econsult, so that means that if you can't use it, you're elderly, English not your first language, you've got learning difficulties... it's not fair. They're doing that whole barrier to protect their time (Clinician 7, GP)

Multiple barriers to safe and effective remote appointments were identified, with these barriers appearing to place certain groups at a disadvantage, particularly patients at an early stage of the

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disease journey. There were reports of greater diagnostic delays, less support at diagnosis, and less chance to build a supportive medical relationship with telemedicine than with face-to-face appointments: *'With new patients it is really important to meet F2F and build up a rapport and confidence'* (Clinician 20, Nurse). These disadvantages were reflected statistically in that the newly diagnosed (<1 year) gave significantly lower ratings (p=0.009) for telemedicine (compared with faceto-face) for relationship-building than those diagnosed >10 years ago.

Telemedicine was also reported by clinicians and patients to disadvantage (Table 3b):

- Patients without English as a first language; those with hearing, cognitive or speech difficulties; or when the accent of the patient or clinician was felt to be difficult to understand.
- Patients with more complex, multi-system, potentially life-threatening diseases, particularly vasculitis or SLE.
- Those in need of urgent care or with MH difficulties.
- Older patients; although the only statistically significant differences in views of telemedicine between age groups was in terms of convenience (a low negative correlation between age and convenience of -0.14, p=0.01).
- Clinicians and patients experiencing difficulties with access to/ability with telemedicine related technology.
- Socio-economically or educationally disadvantaged patients.

Table 3a. Patient types/groups, some of whom expressed a preference for telemedicine

Characteristic	Example quote
Employed	Telemedicine has been great for convenience as someone who works full time. F2F appointments in the NHS are frustrating as they are literally never on time. So have to allocate 2-3 hours for travel and waiting for a 10 min appointment (Ppt 71, Inflammatory arthritis)
Stable, quick check-in	I like the fact that if my condition is stable I don't have to waste 3 hours of my day to tell the consultant that. I have never felt rushed over the phone (Ppt 942, RA)
Experienced	I know my disease and my body. My RA team know that if I say I'm ready for infusion they accept that (Ppt 1237, RA)

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patients	
High	I also feel like it's more convenient as if I am having a bad fatigue day, I can still speak to
fatigue/pain	someone whereas I may have had to cancel an in person slot. (Patient 97, mixed)
or physical	
disability	

Table 3b. Patient and physician characteristics that may make telemedicine less acceptable/effective

Characteristic	Example quote
Undiagnosed	With my initial appointment first postponed and then moved to a phone call it took almost a whole year longer to be diagnosed, a whole year was lost without medication. I think for a first appointment this was terrible (Ppt 989, Inflammatory arthritis)
Newly diagnosed	GP telemedicine was difficult during early diagnosis as everything felt rushed and emotional support was lacking. I found my diagnosis very scary whilst trying to cope with a lot of pain and with very restricted mobility. This element was lost during a telephone appointment whereas at a face to face it would have been very apparent just how difficult I was finding it to cope (1218, RA)
The more complex diseases	I think a lot of us want to do more face to face in particular with something like lupusFor other diseases you just can look at the blood tests and do an adequate job but in something like lupus where blood tests don't always correlate with symptoms and activity you're then going on the only kind of assessment you can do on the telephone which is essentially doing it blind. (Clinician 24, Registrar)
Communication -Accents	Several of the Drs have accents that I find very hard to understand over the phone and as I have a slight hearing loss I find the experience frustrating and uncomfortable (Patient 16, Sjogrens)
Elderly and/or deaf	All telemedicine is hard for the elderly. I am very deaf and even with two hearing aids get stressed talking on the phone about important things (Patient 605, Vasculitis)
Hearing and/or cognitive difficulties	My hearing isn't the best and it's much easier face-to-face. I like to have my wife with me to corroborate and amplify and remind me if forget anything which is difficult on the phone (Patient 1422, lupus)
Less technologically able (clinicians and patients)	My younger colleaguesthey are much better than me at managing online and connection I end up making the mistakesa lot of time wasted in trying to get these online functioning properly as well. (Clinician 23, Consultant)
Socio- economically disadvantaged	In terms of support and living in stressful, over-crowded environmentswon't get what they need or learn easily from a remote consultationneed more support because there's less support at home and they have less background knowledge (Clinician 2, Consultant)

Theme 5: Cost-cutting, clinical need or choice?

Although there was an overall preference for face-to-face appointments, most participants

supported a mixture of face-to-face and telemedicine (Figure 3a). A significantly (p<0.01) higher

proportion of clinicians (70%) than patients (46%) wanted all emergency appointments to be face-to-

face. This disparity may be partially explained by clinicians being very concerned about the

difficulties of diagnosing and treating a flare remotely, including medico-legal considerations, whilst

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patients appreciated the option of telemedicine if feeling too unwell to physically attend. There were no significant differences in patient preference between the mode of telemedicine (phone or video), or between disease groups, in views of telemedicine's acceptability. Clinicians had a strong preference (over twice as many) for telephone appointments over videocalls (Fig 3b). Male clinicians and rheumatology registrars were the most negative clinician sub-groups on the relative merits of telemedicine over face-to-face both quantitively and qualitatively, and rheumatology nurses were the most positive.

Concerns were expressed that telemedicine would be over-used by the NHS and hospital management (Fig3a, 3c) as a cost and time-saving measure rather than in patients' best interests. Clinicians (only 3% of whom felt telemedicine overall was better than face-to-face) and patients had rarely been consulted as to their preferences:

We've got a top-down system where the view on consultation has been taken at a different level than purely a clinical one (Clinician 1, Registrar)

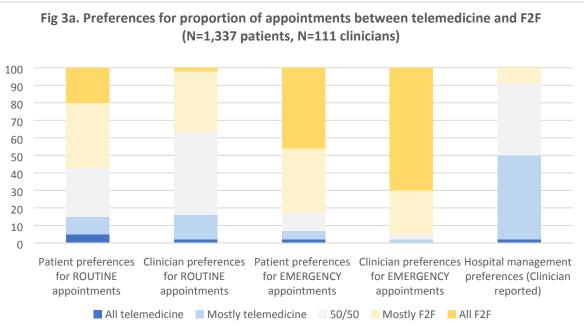


Figure 3. Preferences for mode of appointment delivery

Note: Hospital management preferences are as reported by clinicians

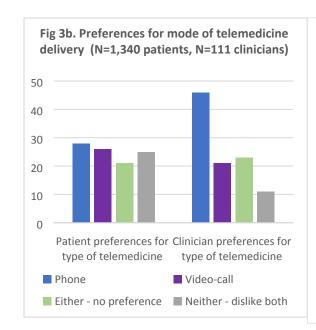


Fig 3c. Example quotes of clinicians' conflicting views with management

I fear that the pandemic will be used to usher in unproven and potentially harmful, but fashionable, changes to practice. There will be some benefits to remote medicine but I fear there hasn't been a serious attempt to think of which clinical situations it is most useful in (Clinician 56, Consultant)

I have a feeling that hospitals will insist on more telemedicine reviews rather than F2F to save money...most patients will hate it (Clinician 65, Nurse)

I feel that management think that equally good care is possible via telemedicine which it is not. The doctor as drug doesn't work as well over the phone (Clinician 108, Consultant)

The importance of assessing individual suitability, for example by pre-consultation questionnaires, and allowing clinician and patient choice, were frequently highlighted:

The rapid digitalisation and use of telemedicine must stay but appropriate patient selection is key, it is perfect for some but disastrous for others (Clinician 40, Consultant)

Discussion

This is the first mixed methods telemedicine study we are aware of that compares the views of rheumatology patients, rheumatology clinicians and GPs. The findings identified a place for telemedicine for certain patients, yet a strong overall preference for face-to-face consultations on the part of both clinicians and patients. Telemedicine was widely perceived to reduce the accuracy of assessments and to impede the building of trusting medical relationships. Consistent with other studies¹⁶, our study indicated that telemedicine was particularly unsuitable for initial rheumatological diagnoses. Contributing factors to the inaccurate assessments and multiple misdiagnoses reported with telemedicine included: no examinations, limited testing, increased

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patient reticence to report symptoms, and increased clinician difficulty in eliciting and identifying unspoken concerns without non-verbal cues. The importance of non-verbal communication in medical relationships has been widely reported^{23,24,25}, including eye-contact²⁶, expressive (to reassure or comfort) touch²⁷ and smiling²⁸. In agreement with other studies²⁹ this lack of non-verbal interaction was found to be particularly detrimental to the eliciting and reporting of mental health concerns, although a minority of our participants (both clinician and patient) found it easier to discuss mental health and sensitive issues remotely. Demonstrations of listening and empathy, identified as of key importance to rheumatology patients in terms of satisfaction and behaviours^{30,31,32}, were perceived by our participants to be limited with telemedicine. This is in contrast to another study reporting very few (<10%) clinicians and patients felt tele-rheumatology had a negative impact on the medical relationship³³.

Our study participants were mainly UK-based (96%), where telemedicine to-date has been reported as often being the substitution of a face-to-face appointment for a telephone-call³⁴, or less frequently a video-call. This may partially explain why our findings were more negative towards telemedicine than studies on medical systems which utilize more visual telemedicine and/or integration of multiple remote methods of assessing and supporting patients^{35,36}. Indeed Leggett at al found a much higher level of diagnostic accuracy and satisfaction with video compared to telephone consultations³⁷. Although interestingly our rheumatology clinicians had a strong preference for telephone over video consultations, possibly influenced by technology constraints and familiarity, and patient preferences were equally divided.

The benefits of digital technology and integrated systems allowing prompt access to health data and communication between patients and clinicians across multiple disciplines and NHS trusts have yet to be realised in many areas. This is a source of inefficiency and frustration currently in the UK³⁸, and addressing it would greatly improve patient empowerment, satisfaction and self-management.

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Greater use of patient-initiated follow-up (PIFU)³⁹ may also reduce existing inefficiencies and suboptimal care from the current dominant model of reviewing rheumatology patients according to timescales rather than disease activity. However, this study has demonstrated that a move towards further PIFU urgently requires improved administrative systems in some rheumatology departments to ensure a prompt, appropriate response to the vast increase in patient-initiated contact. This is currently generating difficulties in managing work-life balance for some clinicians, and endangering and upsetting the many patients who report being unable to access care when required, as also found in studies at earlier stages of the pandemic^{40,41}. Our results indicate that telemedicine may have increased existing barriers to accessing medical support⁴², such as inappropriate telephone triaging, and difficulties navigating administrative and reception systems⁴³. The importance of prompt accessibility in a flare identified in our previous study³⁸ was repeatedly reiterated by senior rheumatologists in this study, as delays can cause permanent organ/joint damage and be fatal in the more severe cases⁹.

We found no differences in preferences for telemedicine between disease groups. However, our participants with inflammatory arthritis had significantly higher confidence in a prompt response from their rheumatology department, higher ratings for medical security, and lower reductions in medical security during the pandemic, than patients with lupus or other rare autoimmune rheumatic diseases, such as Sjogren's. This adds credence to the clinician participants' and authors' concerns that there are inequalities in care between rare and more common rheumatic diseases, and that telemedicine may increase these, and other inequalities. It is particularly important that proactive efforts⁴⁴ through policy ensure telemedicine does not increase existing health inequalities⁴⁵ and further disadvantage groups, such as the elderly and those with communication challenges.

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Aside from the obvious benefits of telemedicine reducing the risks of infection, additional benefits were identified. With a patient population often suffering from pain and fatigue^{31,46,47}, and long

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journeys to hospital, especially for tertiary care, the option of a telemedicine appointment was felt to be equally/more acceptable in some cases. This included for medication reviews or administrative queries, and for stable patients who are knowledgeable, and secure in their pre-existing medical relationship(s). As previous studies also advise, individual choice¹⁶ and careful patient selection for telemedicine will be essential to ensure safety and acceptability, and to alleviate concerns that telemedicine in the NHS is being instituted primarily as a cost and time-saving measure.

It is important to emphasise that our study patients all had autoimmune inflammatory rheumatic diseases and these patients often have more severe and/or unpredictable disease that may be less suited to telemedicine. A limitation of this study is that those with technological, communication or socio-economic barriers to accessing telemedicine are also less likely to participate in an online survey. This study's great strength was in eliciting multiple viewpoints from all sides of patient-clinician relationships, including patients with a variety of rheumatological diseases. This, and combining of qualitative and quantitative data at each stage of the research, allowed for triangulation and helped address threats to validity and reliability. The many studies reporting authors' own patients' high levels of satisfaction should be viewed with some caution due to the power differential and high-risk of social desirability bias. Further strengths and limitations are summarised in Supplementary Data S1, available at *Rheumatology* online.

It is probable that some of the negative perceptions of telemedicine, particularly relating to some access barriers and clinicians having less time, are partly influenced by ongoing Covid-19 pressure and the huge backlog of rheumatology patients requiring appointments, rather than telemedicine itself. Concerns and risks may lessen as remote consultation systems are improved, and the identified benefits for select patients are realised. However, it is unlikely that the serious concerns expressed by our participants regarding assessing complex rheumatological conditions remotely will be fully resolvable. This has implications for the telemedicine ambitions and plans of the UK and

other health systems, and highlights the need for ongoing assessment and mitigation of the clinical

risks associated with a telemedicine strategy.

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