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USABILITY OF A SMARTWATCH FOR ATRIAL FIBRILLATION DETECTION IN OLDER ADULTS AFTER STROKE

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Title. USABILITY OF A SMARTWATCH FOR ATRIAL FIBRILLATION DETECTION IN OLDER ADULTS AFTER STROKE

Short title. Smartwatch AF Detection Post-stroke

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Abstract

Background: Smartwatches can be used for atrial fibrillation (AF) detection, but little is known about how older adults at risk for AF perceive their usability.

Methods: We employed a mixed-methods study design using data from the ongoing Pulsewatch study, a randomized clinical trial (NCT03761394) examining the accuracy of a smartwatch-smartphone app dyad (Samsung/Android) compared to usual care with a patch monitor (Cardea SOLO™ ECG System) for detection of AF among older stroke survivors. To be eligible to participate in PULSEWATCH, participants needed to be at least 50 years of age, have had an ischemic stroke, and have no major contraindications to anticoagulation therapy should AF be detected. After 14 days of use, usability was measured by the Systems Usability Scale (SUS) and investigator-generated questions. Qualitative interviews were conducted, transcribed, and coded via thematic analysis.

Results: Ninety participants in the Pulsewatch trial were randomized to use a smartwatch-smartphone app dyad for 14 days (average age: 65 years, 41% female, 87% White), and 46% found it to be highly usable ($SUS \geq 68$). In quantitative surveys, participants who used an assistive device (e.g. wheelchair) and those with history of anxiety or depression were more likely to report anxiety associated with watch use. In qualitative interviews, study participants reported wanting a streamlined system that was more focused on rhythm monitoring and a smartwatch with a longer battery life. In-person training and support greatly improved their experience, and participants overwhelmingly preferred use of a smartwatch over traditional cardiac monitoring due to its comfort, appearance, and convenience.

Conclusion: Older adults at high risk for AF who were randomized to use a smartwatch-app dyad for AF monitoring over 14 days found it to be usable for AF detection and preferred their

use to the use of a patch monitor. However, participants reported that a simpler device interface and longer smartwatch battery life would increase the system's usability.

Keywords

Atrial fibrillation; smartwatch; Stroke

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Introduction

Atrial fibrillation (AF) is a highly prevalent cardiac rhythm disorder that affects over six million people in the US and over 30 million individuals worldwide, though a large portion of individuals with the condition go undiagnosed due to AF's often asymptomatic and paroxysmal nature.¹⁻³ However, even brief episodes of AF are associated with a greatly increased risk of stroke, and this additional risk is especially magnified in older adults.⁴ Indeed, it is estimated that ischemic stroke is the initial clinical manifestation of about one in five cases of AF.⁵ Furthermore, strokes associated with AF are also more severe than non-AF related ones, and stroke in general contributes to a significant portion of the morbidity and mortality associated with the condition.⁶ Fortunately, treatment options for stroke prevention in AF patients have been shown to be highly effective and safe, which further highlights the importance of timely diagnosis and treatment of the arrhythmia.^{7,8} This is especially important in patients who have already experienced a stroke, as they are a particularly vulnerable population already at higher risk for recurrent stroke, and risk is further exacerbated if they have AF.⁹

Standard of care in patients after a stroke, and especially in patients whose strokes are of undetermined origin, entails routine cardiac rhythm monitoring which most often takes the form of a device with several wires affixed to the patient's chest with adhesive gel.¹⁰ This burdensome modality complicates many activities of daily living, such as exercise or even bathing, and thus suffers from poor patient satisfaction and use.¹¹ In recent years, the a variety of wearable biosensors, and specifically smartwatches, have become available for AF detection. Currently, there are a many consumer-available smartwatches that are cleared by the Food and Drug Administration for using in detection of AF.^{12,13} However, despite tremendous advancements in sensor and computing technologies that have enabled relatively high accuracy of these

smartwatches for AF detection, there has been little consideration of their usability in the older adult population who would benefit most from AF monitoring, particularly for patients who have experienced a stroke. Though smartwatch use in the general population has steadily risen over the last few years, they are by no means ubiquitous, especially among older adults who traditionally have been relatively late adopters of nascent technology.¹⁴ Additionally, the post-stroke population faces an additional host of challenges that may further complicate smartwatch use, such as potential residual neurological deficits or resultant cognitive or psychiatric challenges from their stroke.¹⁵

Relatively few existing mobile and digital health studies are focused on the older adults, and only a portion of them examine device usability. However, advancements in the technological capabilities of wearable sensors have fueled interest in the conversation. The American Heart Association recently published a scientific statement emphasizing the promise of mobile health technologies for preventing cardiovascular disease in older adults, as well as the dire need for a more comprehensive understanding of the barriers that older adults face in using them.¹⁶ Emerging evidence of smartwatch use in older adults has identified potential usability considerations unique to this population, such as accessibility (larger font and icons), intuitive interface design, and potential need for assistance.¹⁷⁻¹⁹ These studies have often been exploratory in nature, and none thus far have focused on patients who have suffered a stroke. We utilized a mixed-methods approach to gain a multifaceted understanding of the usability challenges faced by older adults in using smartwatches for AF detection, which would be crucial for realizing their full potential as a heart rhythm monitoring modality.

Methods

Study design and population

This study uses data from the Pulsewatch study, a randomized controlled trial in which participants were randomized in a 3:1 fashion into intervention and control groups. The intervention group received the Pulsewatch system: a smartwatch-smartphone app dyad capable of AF detection via pulse plethysmography (PPG), while both the control and intervention groups received standard of care ECG patch monitoring (i.e. no smart devices) for a 14-day monitoring period. While the Pulsewatch study consists of a second phase following this 14-day period, because the focus of the present study is on smartwatch usability, we used data only from the intervention group from the first phase of the study (**Figure 1**). Eligible participants were recruited from the neurology and cardiology clinics of a single tertiary care center in central Massachusetts, and were over 50 years old, had an ischemic stroke or transient ischemic attack (TIA) in the last decade, and were willing to use the Pulsewatch system over the course of the study. Exclusion criteria included prior diagnosis of atrial fibrillation, having any major contraindications to anticoagulation therapy, inability to provide informed consent, contraindication for wearing an ECG patch monitor (e.g., sensitivity or allergy to medical adhesives, implantable pacemaker), or a life-threatening arrhythmia requiring immediate analysis and in-patient monitoring.

Study procedures

Eligible patients were identified through the electronic medical record and mailed an invitation letter briefly describing the study and including a study number to call if they had any questions regarding the study or if they wanted to opt out of further contact. These potential participants were then approached at the time of their clinic appointment. If they chose to enroll in the study, they provided informed consent and filled out a baseline study questionnaire

assessing a multitude of sociodemographic and psychosocial domains. The participant was then randomized into either the control or intervention groups. Both groups received standard-of-care cardiac rhythm monitoring in the form of an ECG patch monitor (Cardea SOLO Wireless ECG Patch, Cardiac Insight). In addition, participants randomized to the intervention group received a smartwatch (Samsung Gear S3 or Samsung Galaxy Watch 3) and an accompanying smartphone with the Pulsewatch app installed. This pair of smartwatch and smartphone apps were developed by the study team with significant input from patients who have had a stroke, their at-home caregivers, neurologists and cardiologists. All of these stakeholders provided their perspectives on how best to optimize a study app designed for AF detection to be deployed in older patients who have experienced a stroke. This end-user input was incorporated into the app prior to deployment in the trial, and included considerations ranging from app interface, including adequately large font size and intuitive menu organization, to physical comfort of device wear, such as using easily adjustable magnetic or Velcro watchbands. The final version of the Pulsewatch system consisted of a smartwatch app that near-continuously monitored a participant's pulse for rhythm abnormalities, informing them if potential AF was detected, as well as a phone app that serves as a dashboard for participants to view their data and resources to learn more about AF and strokes. Participants who were randomized to use the Pulsewatch system were asked to consistently wear the watch as much as possible and to engage with the phone app as much as they liked. Research staff called intervention participants on the third and seventh days of the study to encourage watch wear and help troubleshoot any technical challenges that arose. Additionally, participants were provided a comprehensive user guide as well as in-person training at enrollment on general device use. This training involved demonstrating all of the smartphone app's functionalities by explaining each screen in detail and

how to navigate between them. All participants were also shown how to properly wear the smartwatch (specifically the importance of ensure that the watchstrap is tight) and charging the devices.

At the conclusion of the 14-day study period, participants completed a questionnaire similar to the baseline assessment, with additional questions regarding their experience using the Pulsewatch system over the past two weeks. Ten purposively selected participants (to represent a range of demographic factors) were additionally asked to participate in an in-depth interview to garner a more comprehensive understanding of the needs and barriers they faced in their user experience. These interviews were recorded and consisted of two parts. In the first portion of the interview, participants were asked to simulate daily use of the Pulsewatch system, namely putting on and removing the smartwatch, charging the devices, and demonstrating how they have been interacting with the phone app, if at all. The interviewer deemed whether each action was successfully completed, and if yes, whether any assistance was required. The second half followed the format of a semi-structured interview, during which participants are asked about their user experience. Examples of usability domains explored with participants include any barriers or challenges they faced and potential ways to circumvent these barriers, motivations for device use, and user experience compared to traditional cardiac rhythm monitoring modalities.

Study measures

This mixed-methods study consists of both quantitative and qualitative data elements. The primary outcome of the quantitative portion of this study was the System Usability Scale (SUS),²⁰ a validated measure of usability utilized by numerous studies of emerging mobile and digital health tools. Additionally, we generated smartwatch-specific Likert scale questions (5 points: 1 = “strongly disagree”, 2 = “disagree”, 3 = “neutral”, 4 = “agree”, 5 = “strongly agree”)

to better understand facets of use that the more generalized SUS potentially may not cover. These questions were developed through an iterative process by a panel of content experts, including digital health researchers (ED, DDM), biomedical engineers (DH, KC), a cardiac electrophysiologist (DDM), and a gerontologist (JSS). The final iteration of questions addressed the participants' perceived ease of use of both the phone and watch, their feeling connected to healthcare providers, any potential anxiety as a result of using the system, and their general use experience. Participants' medical characteristics were abstracted from the EMR by trained research staff. Cognitive status was assessed with the Montreal Cognitive Assessment (MoCA),²¹ depressive symptoms were examined using the Patient Health Questionnaire (PHQ-9),²² and anxiety symptoms were assessed with the Generalized Anxiety Disorder (GAD-7)²³ instruments.

Analytic approach

Descriptive statistics were calculated to summarize participant-level factors for both the overall cohort as well as the ten participants who also completed in-depth interviews. Linear regression models were used to examine associations between individual participant-level characteristics and SUS score. Independent variables in the models were chosen based on prior literature, as well as other factors that are likely to be associated with usability. We also conducted secondary analyses using logistic regression models with SUS as a binary outcome (dichotomizing with a previously validated cutoff of 68 for high vs low usability²⁰). The investigator-generated Likert scale questions are presented as the proportion of participants who responded with a 4 or 5 (i.e., "agree" or "strongly agree"). Additionally, each question is used in an ordinal logistic regression model with the full "strongly disagree" to "strongly agree" scale as the dependent variable, with the same predictor variables as the models described above.

Interview recordings were transcribed and reviewed for quality control. Qualitative coding via directed content analysis was completed by two independent researchers (ED, MCA) and any differences were resolved through discussion and mutual agreement. Themes from the analysis originate from concepts outlined in previous literature, and the coding guide was developed after an initial review of transcriptions through an iterative process, and was refined through discussion between research team members (ED, KM). The quantitative and qualitative results were triangulated, and the qualitative outcomes were used to contextualize the quantitative findings from the study.

Results

Quantitative results

A total of 90 participants were randomized to receive the Pulsewatch system. The average age of the cohort was 65 years old (SD 9), 41% were women, and most were white (87%). A large proportion of participants were of high socioeconomic status, with 52% having a college degree and 33% who had an annual income of over \$100,000. Thirty percent were cognitively impaired, though there was relatively low burden of depressive and anxiety symptoms in the sample. Our participants were highly engaged with technology, especially given the age group, with 83% of participants owning a smartphone, 25% already owning a smartwatch, and two-thirds of participants using apps on their devices on a daily basis (**Table 1**).

With regards to the SUS, 39% of participants deemed the system highly usable ($SUS \geq 68$). More than half of participants either “agreed” or “strongly agreed” with both the watch and phone app being easy to use, and 42% endorsed that using the system made them feel more connected to their healthcare providers. About one-eighth of participants experienced anxiety

from using the devices, and more than half of the cohort agreed that they enjoyed their experience using the devices (**Table 2**).

Age was associated with increased likelihood to experience anxiety or worry with use of the system, and every additional year is associated with approximately 6 % odds (OR 1.06, 95% CI: 1.01 to 1.11) of moving up 1 point on a 5-point Likert scale (with 1 being “strongly disagree” with the statement that using the system caused anxiety, and 5 being “strongly agree”). Participants who reported use an assistive device (defined as cane, walker, wheelchair), or have anxiety or depression at baseline are more likely to report anxiety or worry with use of the system (**Table 3**). No other demographic, medical history, psychosocial, or technologic familiarity variables investigated were associated with either the SUS or any investigator-generated usability questions.

Qualitative results

Characteristics of the sub-cohort of participants who were selected to do an in-depth interview are presented in **Table 4**. Interview lengths ranged from 16 to 51 minutes. , In the demonstration portion of the interview, all participants were successful in independently placing the watch on their wrists and adequately tightening the watchstrap, removing the watch, and charging devices with no external assistance. Eight of the ten participants engaged with the accompanying phone app to varying degrees, and all 8 were able to successfully navigate through the app to visualize their own heart rate and rhythm data, though multiple participants beyond those selected for the qualitative portion noted to study staff that they rarely engaged with the phone app.

Device training and resources

Participants indicated that the training they received initially upon receiving the devices from study staff was instrumental to successfully using the devices; this was particularly true for

those who do not own smartwatches. Demonstration of use of the watch and phone apps appeared to remove a major usability barrier by familiarizing the participant prior to individual use. This live session with research staff was perceived as far superior to the detailed written instructions with images that were also provided. Some participants preferred a more brief, bulleted list in lieu of the detailed instructions provided, and those who did find the detailed written instructions helpful mentioned that they generally used the document as a back-up reference, and did not often utilize the guide.

Training made it way easier, because she [research staff] explained everything in full, and I don't know, I probably wouldn't have done it if she didn't tell me exactly what to expect, and what I was going to do, you know? The training is a necessity. Absolutely. Especially the older people, because they're going to be so confused [otherwise].

- Female, age 53, owns a smartphone

This preference for support in the form of dedicated personnel extended beyond the initial encounter, and participants found the ability to reach study staff any time they encountered any issues to be tremendously useful. This was true even for participants who did not utilize the study phone number for troubleshooting, as they reported that just the knowledge that they could contact study personnel for assistance if needed contributed positively to their experience.

Most participants had family members at home who they perceived to be more technologically adept than they were, and were able to help them with devices if needed, though it was often not necessary due to the constant availability of study staff and the perceived ease of use of the system. This ease of use was attributed to the passivity of monitoring, and the relatively low amount of active engagement asked of participants.

I have people that would be more technically advanced than me, if there was something I needed. But I mean, I just saw that I was supposed to turn it on [the watch], and have that icon come up and see that it [the phone app] was open once a day, so I could see that my heart rate and everything else. I had nothing else that I needed to do.

- Female, age 61, owns a smartphone

Streamlining rhythm monitoring and minimizing patient burden

While participants generally indicated that the Pulsewatch system was easy to use, many cited streamlining the system to be a crucial factor for any digital health tools to be deployed in this population. Although newer generation smart devices are being embedded with an increasingly wider array of functionalities, from texting and calling to mobile payment, participants indicated that these additional functions only served to confound the user experience and detracted from the main goal of cardiac rhythm monitoring:

There were just too many other things that you could do ... the next thing you know, you're on the weather [app] – “wait a minute, I don't want that.” So, it would be better if they had a [system] that was not as complete as one of these, just having the ability to do the heart recording and monitoring, but not having all these other functions. Without having all the extraneous applications and stuff on there, it would make it so much easier to function, to focus on.

- Male, age 61, owns a smartphone

While streamlining and focusing the functionalities of the watch was a recurrent theme, several participants also noted that the general health monitoring provided by smartwatches, such as physical activity or sleep metrics, are potentially helpful to visualize as well, and may be

acceptable to include in a cardiac rhythm monitoring system. However, among all participants, including those who noted the potential helpfulness of these additional functionalities, there was still particular emphasis on the necessity of a passive system that does not require extensive active engagement on the part of the patient:

If your goal is to have an app that is all-encompassing, then a step tracker and some of those other features would be great. But if it's really to try to replace an invasive loop monitor, it needs to be super passive. It needs to be a "set it and forget it".

- Male, age 53, owns smartphone and smartwatch

A passive system would be perfect. Because I wouldn't have to [actively] record anything myself, like everything is there. So, it would be pretty easy and ... I do not want to touch it.

- Female, age 51, does not own smartphone or watch

Compared to traditional monitoring modalities

Most participants interviewed had prior experience with either a Holter monitor, external event monitor, or implantable loop recorder. Participants overwhelmingly indicated a strong preference for the Pulsewatch system if given the choice of cardiac monitors, provided that the watch could attain the same diagnostic accuracy as clinical gold-standard monitors. The appearance of the watch was a factor for some participants who were dissatisfied with the obtrusive and bulky appearance of Holter monitors. Comfort was also the most commonly cited reason, specifically given the various ways that external cardiac monitoring significantly impacts quality of life and daily activities.

This [watch] would be better. Well, it's not as intrusive. I mean, I'm wearing this big thing [event monitor] and every time I turn over in bed, it'll pull on me and wake me up, so it's very tough to deal with when you're wearing it all the time. I had to cover it [event monitor] up to take a shower and stuff, so this [watch] is great, it really was good.

- Male, age 87, owns a smartphone

Oh absolutely. I could go out [with the watch] and no one would notice it. There wouldn't be wires hanging out of my shirt, people staring at me like what the hell is wrong with her, you know?

- Female, age 51, owns smartphone

An additional advantage of the Pulsewatch system that some participants noted over traditional monitoring modalities is the ability to visualize their own data in near real time on the accompanying phone app. One participant reported that her heart rate dropped to 37 beats per minute while wearing the smartwatch, and while not AF related, she was able to monitor her heart rate displayed on the watch in real time in order to make a conscious and informed decision on whether or not to seek urgent medical care.

Sense of security

Participants viewed wearing the smartwatch as providing them with a sense of security and peace of mind. The Pulsewatch system was often cited as a “safety net” by participants, especially by those who have had acute cardiac events. This aspect of the system was the biggest incentive to use a commercial wearable device for them. Conversely, several participants experienced anxiety while using the Pulsewatch system attributable to the stress of potentially

missing AF episodes during times when they were not wearing the smartwatch, for example, while it is charging or when they were in the shower. One participant cited the waterproof feature of the watch to be useful specifically for the purpose of being able to wear it in the shower, to minimize the duration of non-wear. Participants recognized the value of continuous AF monitoring, and any gaps in this monitoring was perceived as a major issue. This was especially salient given that due to the near-continuous nature of pulse monitoring by the smartwatch (and therefore battery expenditure), it ran out of power in about 6 to 8 hours, and needed to be charged multiple times a day. These periods of non-wear therefore conferred anxiety and was a major source of frustration for participants:

Battery life was definitely a big issue. It was on the charger more than my hand! But I was more relaxed wearing the watch actually, because of my heart thing. Like maybe it [the watch] would be able to show something. I felt more comfortable with it on, when it had power. Because if it's not going to be on, then what's the point?

- Female, age 73, does not own smartphone or watch

Health as a motivator for system use

The major motivation for participants to use a wearable system for heart rhythm monitoring was to maintain their health, whether it was intrinsically motivated or to generate data for their healthcare providers. Having this information contributed to the sense of security described. One participant cited the Pulsewatch system as helping normalize her life after her stroke, as the constant feedback on her heart rate and rhythm gradually restored her confidence to engage in exertional daily tasks, such as gardening, that she felt anxious about doing

immediately following the stroke. Additionally, some participants noted the novelty of a smartwatch for rhythm monitoring may help motivate them to use the device:

Just having it monitoring my heart and noticing if anything goes wrong, or if everything is just normal. And also, the novelty factor, having new stuff to play with.

- Male, age 61, owns a smartphone

Integrating quantitative and qualitative results

Quantitative and qualitative data elements in the study were integrated based on triangulation protocol to describe whether data elements were convergent (where findings from both methods agree), complementary (where results from qualitative and quantitative methods enhance understanding of a particular phenomenon), or divergent (where findings appear to contradict one another).²⁴ We observed overall agreement, and the qualitative component of this study contextualized the quantitative observations (**Table 5**).

Discussion

In this usability study of smartwatch-based AF detection in older adults after stroke, we observed that participants found the Pulsewatch system be generally easy to use due to its passivity, though a more streamlined approach focused only on cardiac rhythm monitoring may have been preferable to a multi-functional commercial smartwatch. A small portion of participants experienced anxiety from using the system. Those who are older, use assistive device or report anxiety or depression at baseline endorsed higher agreement with the system causing them anxiety, suggesting that they may benefit from more traditional methods of AF detection. Finally, participants overwhelmingly preferred smartwatches as a heart monitoring

modality to traditional clinical monitors. Their preference stemmed from logistical reasons related to wear factors such as comfort, adherence, and convenience.

Contextualizing usability score

Usability of the Pulsewatch system was rated very similarly to other commercially available smartwatches, though there is limited research conducted in older patients with stroke. A usability study of 7 different models of commercially available smartwatches, including the Samsung Gear S3 used in the present study, found that average SUS scores of the devices ranged from 61.4 to 67.6 (versus 65.1 in this study), though notably, the study recruited existing smartwatch users, and only one of the nearly 400 participants was over the age of 60.²⁵ Additionally, a recent mixed-methods study of eight older adults (mean age = 62) reported a median SUS score of 60 for using wrist-based activity tracker.²⁶ The stability of SUS score ratings of smartwatches across drastically different populations may indicate that though older adults may be initially less willing to embrace emerging technologies, their actual use of these devices may not differ significantly from the general population. Additionally, even the specific challenges associated with a stroke do not necessarily preclude the use of smartwatches for cardiac rhythm monitoring. This is further exemplified by our finding that no demographic, medical history, psychosocial, or technological familiarity factors were significantly associated with SUS score. The majority of participants indicated that the system was easy to use, but characteristics such as education, cognitive impairment, or previous device ownership were interestingly not associated with any usability domains examined. This suggests that a simple and passive smartwatch-based system for cardiac monitoring may be accessible for many older patients after stroke, irrespective of many individual-level factors.

Facilitators of smartwatch usability

However, though the Pulsewatch system was generally rated as fairly usable, there are many challenges unique to this population that impacts their user experience. An important and recurring theme was the need for streamlining the system to focus primarily on measuring health metrics without convoluting the process with extraneous functionalities. While manufacturers in the industry sector race to integrate new functionalities into successive generations of devices, it would be prudent to more thoroughly consider the needs of older populations for whom many of these health features are designed for, and devote more resources to streamlining the user interfaces of these devices to be more intuitive for older individuals. Additionally, passivity was deemed crucial to ensuring that the devices were easy to use for participants. A usability study of Fitbit devices among adults over the age of 65 found that when asked to set up their own Fitbit accounts and entering their own data through the app, participants were reluctant to use the devices and deemed the process to be not worth the effort.²⁷ Decreasing participant burden by minimizing their active role appears to be an important factor in smartwatch usability for older adults. This is further corroborated by other studies indicating that older adults found a wrist-based activity tracker to be simple to use specifically due to the lack of interaction required on their part.²⁶ Finally, dedicated training and support personnel appears to drastically improve the user experience for the system among participants. This initial training was perceived as especially important, and not something that could be replaced by written instructions. The interpersonal nature of this interaction is likely key to successful use, as older adults have been shown to perceive wearable devices to be less complex when they are able to observe others using it.²⁷

One issue noted by several participants was the short battery life on the smartwatches. Previous research suggests that a majority of older adults consider long battery life to be essential for ensuring long-term use of wearable devices, indicating that one week would be ideal and that having to charge the device daily was unacceptable.²⁶ While the Pulsewatch cohort had lower expectations of the devices with respect to being able to last a week without having to charge, limited battery life was nonetheless a common complaint among participants. For most participants, having to charge the devices frequently seemed to be a mere annoyance, but several experienced anxiety as a result of having gaps in their cardiac rhythm monitoring. This impacts quality of life for potential smartwatch users, and should be taken into consideration by manufacturers of these devices. A potential method to prolong smartwatch battery life may be to reduce its functionalities to the core components necessary for rhythm monitoring, and thus eliminating additional demands for processing power and battery expenditure. Furthermore, this would be in line with the ease-of-use considerations mentioned above in streamlining the functionalities and interfaces of these devices, thereby facilitating use.

Strengths and limitations

Our study has multiple strengths. It fills a critical research gap in studying older adults' perceptions of and willingness to use a smartwatch-based cardiac rhythm monitoring system when a majority of mobile and digital health studies focus on younger populations. Additionally, our mixed-methods approach enabled a comprehensive and multi-faceted investigation, allowing for a synergistic understanding of crucial considerations with regards to usability of smartwatches in this population. This study also has several weaknesses. The study cohort is homogenous with regards to race and ethnicity and had experience with technology use at

baseline, so the findings of this study may not be generalizable across different populations. Generalizability may be further limited by the use of one specific software and hardware combination in this study. Additionally, the present study's sample size was small and thus may not be powered to identify group differences with regards to usability or other factors.

Conclusions

Older adults with stroke generally found a smartwatch-based cardiac rhythm monitoring system to be usable, though a more streamlined system focused purely on passive cardiac monitoring, with longer battery life would likely be more ideal for this population. However, despite these challenges, older adults highly preferred the smartwatch system compared to traditional cardiac monitoring due to its comfort, appearance, and convenience.

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a member of the Heart Rhythm Society and the editor-in-chief of Cardiovascular Digital Health Journal. The other authors have no disclosures.

Authorship

All authors attest they meet the current ICMJE criteria for authorship.

Patient Consent

All patients provided written informed consent.

Ethics Statement

The authors designed the study, gathered and analyzed the data according to the Helsinki Declaration guidelines on human research.

The research protocol used in this study was reviewed and approved by the institutional review board.

Disclaimer

Given his role as Editor-in-Chief, Dr. David McManus had no involvement in the peer review of this article and has no access to information regarding its peer review. Given his role as Associate Editor, Dr. Ki Chon had no involvement in the peer review of this article and has no access to information regarding its peer review. Full responsibility of the editorial process for this article was delegated to David Duncker, MD

Table 1 – Baseline Characteristics of Participants

Demographics	n = 90
Age, mean, yrs (Standard Deviation [SD])	65.1 (9.3)
Female	37 (41%)
Race	
White	78 (87%)
More than one race	6 (7%)
Black	1 (1%)
Asian/Pacific Islander	1 (1%)
Other	4 (4%)
Non-Hispanic ethnicity	87 (97%)
Married/living as married	61 (69%)
Education	
Less than high school	4 (5%)
High school degree or equivalent	38 (43%)
College degree	28 (32%)
Post-graduate degree	18 (20%)
Income	
< \$50,000	29 (35%)
\$50,000 - \$99,999	27 (33%)
≥ \$100,000	27 (33%)
Medical history	
History of ischemic stroke	71 (79%)
History of TIA	26 (29%)
Congestive heart failure	6 (7%)
Cardiac arrhythmias	12 (13%)
Valvular disease	9 (10%)
Hypertension	70 (78%)
Chronic pulmonary disease	7 (8%)
Diabetes	25 (28%)
Vascular disease	24 (27%)
Renal disease	4 (4%)
Prior major bleed	5 (6%)
Prior MI	16 (18%)
Hyperlipidemia	77 (86%)
Sleep Apnea	25 (28%)
Percutaneous Coronary Intervention	11 (12%)
Residual neurological deficit	28 (31%)
Medication use	
Antiarrhythmic	2 (2%)
Beta blocker	40 (44%)
Calcium channel blocker	62 (69%)
Anticoagulant	11 (12%)
Antihypertensive	51 (57%)
Antiplatelet	79 (88%)

Statin	82 (91%)
Vital signs	
BMI, mean (SD)	32.0 (21.0)
Systolic BP, mean (SD)	131.4 (16.7)
Diastolic BP, mean (SD)	76.0 (8.6)
Heart Rate, mean (SD)	73.1 (14.7)
Psychosocial variables	
Alcohol use	7 (8%)
Cognitive impairment, MoCA < 23 for in-person, < 17 for phone	26 (30%)
Vision impairment	48 (54%)
Hearing impairment	26 (29%)
Depressive symptoms	
None (0 – 4)	49 (54%)
Mild (5 – 9)	27 (30%)
Moderate (10 – 14)	8 (9%)
Moderately severe (15 – 19)	3 (3%)
Severe (> 20)	3 (3%)
Anxiety symptoms	
None/minimal (0 – 4)	62 (69%)
Mild (5 – 9)	16 (18%)
Moderate (10 – 14)	7 (8%)
Severe (>15)	5 (6%)
Technology engagement	
Device ownership	
Tablet	60 (67%)
Smartphone	74 (83%)
Smartwatch	22 (25%)
Basic cellphone (SMS enabled)	30 (34%)
App use frequency (excluding call/text)	
Daily	54 (68%)
A few days a week	12 (15%)
At least once a week	5 (6%)
Less than once a week	2 (3%)
Once a month	3 (4%)
Never	4 (5%)

Table 2 – Participant perceptions of the Pulsewatch system

Usability domain	Percent of participants in agreement (n = 90)
Easy to use (smartwatch)	63.6%
Easy to use (smartphone app)	52.3%
More connected to healthcare providers	42.5%
Anxiety or worry from use	12.5%
Overall enjoyed user experience	51.1%

Table 3. Factors associated with smartwatch causing anxiety

Characteristics	Unadjusted OR
Age	1.05 (95% CI: 1.01 to 1.11)
Use of assistive device*	3.34 (95% CI: 1.12 to 9.96)
Baseline depression symptoms	2.31 (95% CI: 1.01 to 5.25)
Baseline anxiety symptoms	3.85 (95% CI: 1.56 to 9.48)

* Assistive device defined as cane, walker, and wheelchair.

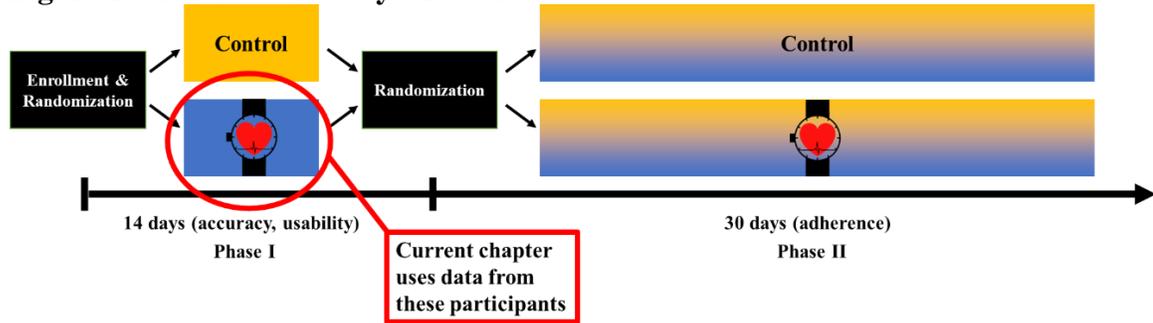
Table 4 – Characteristics of participants who completed an in-depth interview

Demographics	n = 10
Age, mean (SD)	62.8 (11.6)
Female	7 (70%)
Race	
White	6 (60%)
Black	1 (10%)
American Indian/Alaska Native	1 (10%)
More than one race	1 (10%)
Other	1 (10%)
Married/living as married	4 (40%)
Education	
Less than high school	1 (10%)
High school degree	5 (50%)
College degree	1 (10%)
Post graduate degree	3 (30%)
Income*	
Less than \$50,000	5 (56%)
\$50,000 - \$99,999	2 (22%)
More than \$100,000	2 (22%)
Device ownership	
Tablet computer	8 (80%)
Smartphone	8 (80%)
Smartwatch	2 (20%)
Basic cell phone	4 (40%)

*One participant chose not to disclose income (n = 9)

Table 5 – Triangulation of quantitative and qualitative usability results

Outcome of interest	Quantitative results	Qualitative results	Triangulation
Ease of use	63.7% of participants found watch easy to use 56.5% found the phone app to be easy to use	- The devices were simple to use because there was little interaction required – passivity is an important facet of making the devices easier to use for older adults - Training and support for dedicated personnel greatly enhanced user experience and made the system easier to use for participants	Convergent / Complementary
Anxiety from use	10.2% of participants reported experiencing anxiety or worry from using the system	- Anxiety from use stemmed from not being monitored continuously due to having to take off the smartwatch for charging or other reasons - Extending the battery life of the smartwatches would have severely curb feelings of anxiety in participants	Complementary

Figure 1 – Pulsewatch study structure

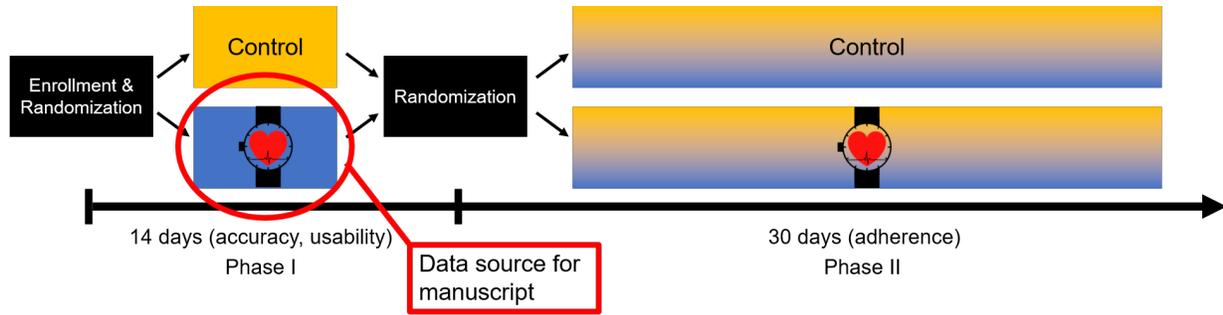
Footnote: data from participants in the Phase I intervention group of the Pulsewatch trial

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Key Findings

- Older stroke survivors find a smartwatch to be generally usable for detecting atrial fibrillation
- Some patients may experience anxiety from using a smartwatch for rhythm monitoring, especially older patients, those who need assistive devices, or have mood comorbidities at baseline.
- Further developments in industry are necessary to streamline functionalities and improve battery life to fully realize the potential of smartwatches for atrial fibrillation detection.

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