Peer-reviewed publications demonstrating the clinical validity and utility of the Zio system
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Diagnostic Yield of Extended Cardiac Patch Monitoring in Patients with Stroke or TIA. *Frontiers in Neurology*, 2014


Ambulatory Cardiac Monitoring for Discharged Emergency Department Patients with Possible Cardiac Arrhythmias. *Western Journal of Emergency Medicine*, 2014


Verba, S.D., Jensen, B.T. and Lynn, J.S.  *Electrocardiographic Responses to Deer Hunting in Men and Women.*  *Wilderness & Environmental Medicine,* 2016

**Keywords:** Arrhythmia detection during exercise

Deer hunting includes various stimuli resulting in augmented sympathetic activity, increased heart rate (HR) response, and rhythm changes. Collectively, these superimposed stresses may increase an individual’s risk for cardiovascular events. This study evaluates HR and rhythm responses in multiple phases of deer hunting in men and women with and without cardiovascular disease (CVD).

- Nineteen participants, 6 female, age 38.3 ± 13.8 years (mean ± SD) with body mass index 29.2 ± 6.9 kg/m² followed their normal hunting routine.
- Three hunters recorded HR ≥85% of their age-predicted heart rate maximum (HRmax) for 1 to 2 minutes.
- Arrhythmias were detected in both participants with CVD and in 8 without CVD: premature atrial, junctional, and ventricular complexes
- Fifteen of 19 hunters experienced “buck fever” (acute extreme excitation), with 7 reaching ≥85% HRmax for up to 1 minute
- The unobtrusive profile of the device resulted in high subject compliance and device adherence during all phases of the hunt. The HRs and ECG recordings had good signal quality for analysis.

Men and women with and without CVD recorded substantial increases in HR and clinically relevant arrhythmias while deer hunting.

Steinhubl, S., et al.  *Rationale and design of a home-based trial using wearable sensors to detect asymptomatic atrial fibrillation in a targeted population: The mHealth Screening To Prevent Strokes (mSToPS) trial.*  *American Heart Journal,* 2016

**Keywords:** Atrial Fibrillation, Silent AFib Screening

Researchers at the Scripps Translational Science Institute (STSI) have launched a home-based clinical trial using the ZIO Service to identify patients with asymptomatic atrial fibrillation (AFib).

- The mSToPS clinical trial aims to determine whether screening select individuals in their homes using wearable sensor technology can detect asymptomatic AFib more efficiently than routine care, such as primary care visits.
- To conduct the study, STSI has teamed with iRhythm, Aetna’s Innovation Labs and Healthagen Outcomes units, Janssen Pharmaceuticals and Amiigo consumer heart rate tracker.
- 2,100 active monitoring participants will be compared to 4,000 usual care beneficiaries.
- Participants will undergo continuous single-lead ECG monitoring using the ZIO XT Patch for the first two weeks and last two weeks of the four-month monitoring period.

**Keywords:** Drug Trials, Heart Failure and Arrhythmia Detection

This publication is a methods paper describing how the clinical trial for Gilead’s investigational drug Eleclazine will progress.

- Hypertrophic Cardiomyopathy (HCM) is complex and not well understood. As HCM progresses, there are a number of potentially serious health consequences that occur, including diastolic heart failure, microvascular dysfunction, atrial fibrillation (AFib) and sudden cardiac death.

- Extended cardiac monitoring with the ZIO Service allows researchers to capture additional relevant information. This may help better determine the pathophysiology of HCM and advance the development of investigational treatment options to address a significant unmet medical need.


**Keywords:** AFib Burden and Cognition, Atrial Fibrillation

Study results showed an association between a high burden of atrial fibrillation (AFib) and lower cognitive function. Previous studies have shown a relationship between AFib, cognitive decline and increased risk of dementia. However, this study demonstrates a correlation between high AFib burden — the percent of time a person has AFib — and cognition.

- The study was based on 325 participants from the Atherosclerosis Risk in Communities (ARIC) Study who wore the ZIO Patch.

- Compared with participants who did not have AFib, participants with AFib burden of 100% (persistent AFib) had lower Animal Naming (AN), Trail Making Test part B, and Digit Span Backwards (DSB) scores. These are standard cognitive assessment tests.

- By contrast, participants with an AFib burden of 1% to 6% did not have lower cognitive test scores than those without AFib.
Kaiser researchers examined 128,401 episodes of monitoring between October 2011 and 2013 using iRhythm’s ZIO Service for which the average monitor wear time was nearly 10 days and more than one quarter were worn for 14 days.

- 18.3% of recordings had at least one episode of non-sustained ventricular tachycardia (NSVT), 0.2% with sustained VT, 1.4% with a sinus pause >3 seconds (SP), 0.4% with a pause during atrial fibrillation >5 seconds (AFP), and 1.2% with high-grade heart block (HGHB)
- Median time to first arrhythmia: 74 hours for NSVT, 22 hours for sustained VT, 22 hours for SP, 31 hours for AFP, and 40 hours for HGHB.
- A significant percentage of potentially high-risk arrhythmias were not identified within 48-hours of ambulatory ECG monitoring. Longer-term continuous ambulatory ECG monitoring provides incremental detection of these potentially clinically relevant events.

Adhesive ECG patch devices are becoming the standard for detecting arrhythmias in the outpatient setting when short to medium term monitoring is indicated. These cardiac devices and related digital mobile health technologies are reshaping the clinician-patient interface with important implications for future healthcare delivery.

- Studies highlight the challenges in diagnosing atrial fibrillation (AFib) with conventional monitoring even in relatively high arrhythmia burden patients with paroxysms. Studies support the use of prolonged ECG monitoring in most patients suspected to have atrial arrhythmia(s) and/or neurologic symptoms suggestive of impending or ongoing TIA or stroke.
- Prolonged ECG monitoring studies have revealed that AFib remains vastly under-diagnosed and that duration of cardiac monitoring following acute ischemic stroke should be extended beyond 24-48 hours (Schuchert et al., 1999; Tayal et al., 2008; Elijovich et al., 2009).
- In another study of 56 patients with cryptogenic TIA or stroke, AFib was diagnosed after a median of 7 days (Tayal et al., 2008).
**Keywords:** Sub-clinical Atrial Fibrillation Detection in High Risk Population, Silent AFib

Prospective study of 75 male patients screened using ZIO Patch detected atrial fibrillation (AFib) and atrial tachycardia (AT) in 11% of asymptomatic patients (silent AFib) with known risk factors.

- Inclusion criteria were age ≥55 years and ≥2 of the following risk factors: coronary disease, heart failure, hypertension, diabetes, sleep apnea. Patients were excluded with prior AFib, stroke, transient ischemic attack, implantable pacemaker or defibrillator, or palpitations or syncope in the prior year.
- AFib was detected in 4 subjects (5.3%; mean AFib burden 28%).
- AT ≥60 seconds was present in 5 subjects (6.7%).
- The combined diagnostic yield of sustained AT/AFib was 11%.
- Found a high prevalence of asymptomatic AT and frequent supraventricular ectopic complexes, which may be relevant to development of AFib or stroke.


**Keywords:** Diagnostic Odyssey, Failure of Holter to Diagnosis Arrhythmias

This is a non-ZIO study. Claims analysis performed using a 5% random sample of Medicare beneficiaries’ claims from Fee-for-Service Standard Analytic Files (SAF). The analysis was limited to patients with full benefits for 1 year prior and 2 years post the index Holter event, with no prior arrhythmia or Holter.

- Clinicians were unable to rule-in or rule-out arrhythmias in 11.1% of the claims evaluated, even after repeated Holter monitoring.
- In spite of this failure, there was a total allowed charge of more than $45 million, which calculates to more than $23,000 per involved patient.
- When extrapolated over the entire Medicare Fee For Service population, this category was estimated to have cost more than $900 million over the 2-year study period.

**Tung, C., Turakhia, M., and Lansberg, M.** Diagnostic Yield of Extended Cardiac Patch Monitoring in Patients with Stroke or TIA. *Frontiers in Neurology*, 2014
Keywords: ZIO Patch + Stroke/TIA Patients

Retrospective study of 1,171 patients monitored using the ZIO Service between January 2012 and June 2013 with an indication of TIA or stroke (excluding cryptogenic).

- Patch monitoring had high patient compliance (median wear time 13.0 days) and analyzable time (98.7%).
  - Demonstrates that the cardiac patch, a novel device for detection of atrial fibrillation (AFib) and other cardiac arrhythmias, is well tolerated by stroke and TIA patients.

- AFib was present in 5.0% of reports.
  - The mean AFib burden was 12.7%, demonstrating the transient nature of the arrhythmia which can complicate detection.

- The high rate of SVT detection (70%) in this patient population with a history of stroke or TIA is noteworthy as it may be a precursor to AFib.


Keywords: Arrhythmia Detection + Long Term Continuous ECG Monitoring

Researchers reviewed data from 524 consecutive patients referred to an academic electrophysiology practice and prescribed a ZIO Patch. Patients were instructed to wear the device for up to 14 days and to activate a trigger button on the device when they experienced symptoms.

- Overall, 99% of patients had some recorded arrhythmia, which included ectopy.

- The most clinically significant arrhythmias were atrial fibrillation/flutter (AFib) in 105 patients (20%), and non-sustained ventricular tachycardia in 79 patients (15%).

- Over one-third of initial arrhythmias were recorded after 48 hours.

- The most common rhythm associated with patient triggered symptoms was normal sinus (50%).

- The majority of AFib episodes (62%) were asymptomatic.

- Long-term ECG monitoring detected arrhythmias in all subjects, and a large percentage were detected after 48 hours. Patient-reported symptoms did not correlate with arrhythmias, including AF, in half of all symptom recordings.

Keywords: ZIO Patch in ED Patients

Retrospective study of 174 adult ED patients with symptoms of possible cardiac arrhythmia who were discharged with a ZIO Patch from one of 3 academic EDs in the US. Study aimed to determine the diagnostic yield of the ZIO Service and the value of prolonged monitoring of these patients.

- The ZIO Service had a higher diagnostic yield of 63% in low-risk patients discharged from the ED, compared to 15% found with 24-48-hour Holter monitoring in previous studies.
- 53% of patients with symptoms, as noted by depressing the ZIO Patch event button, did not have an arrhythmia present at the time. This symptom-rhythm correlation is helpful when “Ruling Out” the presence of arrhythmia when symptoms are noted.
- The median time to the first triggered arrhythmia for potentially serious arrhythmias (ventricular tachycardia and pauses >3 seconds) was 3.1 and 4.2 days, outside of the detection window of traditional Holter monitoring.
- Ease of use with the ZIO Patch was demonstrated by 100% of participants successfully returning the device.
- The ZIO Service is clinically useful in an ED setting as it provides relatively prompt diagnoses of both normal sinus rhythm in symptomatic patients as well as serious asymptomatic arrhythmias in others.


Keywords: Arrhythmia Detection with ZIO Patch Compared to Holter

Prospective study of 146 consecutive patients referred for evaluation of cardiac arrhythmia who underwent simultaneous ambulatory ECG recording with the ZIO Patch and 24-hour Holter monitor.

- The ZIO Service detected 57% more arrhythmias: 96 arrhythmia events by the ZIO Service as compared to 61 arrhythmia events by the Holter monitor (p<0.001).
- 90% of the time, referring physicians reported the ZIO Service aided in a definitive diagnosis compared to 64% for Holter monitoring.
- 81% of patients preferred ZIO Patch over Holter monitoring, which contributes to a longer wear time and improved arrhythmia detection.

Keywords: Premature Ventricular Contraction, ARVD/C Diagnosis

First study to examine the 7-day variability in PVC frequency in patients with Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C). 40 patients received ZIO Patches from the Johns Hopkins ARVD/C registry.

- Patches were prescribed for 7 days and worn an average of 6.6 days.
- Substantial statistical variability in PVC counts was found between 24-hour periods. PVC burden was shown to be present in 76% of patients in this study. If only a single 24 hour Holter is applied, the 24-hour PVC count may be above or below the 500 PVC/24-hour threshold used for ARVD/C diagnosis.
- However, in patients already diagnosed with ARVD/C, the degree to which PVC variability is likely to impact clinical practice is not as well-known and requires more research.


Keywords: Syncope, Diagnostic ECG Monitoring

Syncope has a broad range of causes and pursuit of a correct diagnosis can be tedious and expensive. Cardiogenic syncope is the most common etiology in the critical care setting; prognosis and risk of cardiovascular mortality is significantly higher compared to other forms of syncope.

- The gold standard in the evaluation for syncope after the initial workup is the documentation of reproducible symptoms, often done either with prolonged ECG monitoring.
- Holter monitors are the least sensitive ECG monitoring technique. The ZIO Patch has been shown to detect more arrhythmias and is less cumbersome to wear than traditional Holter monitors. It should be noted that the yield of Holter monitoring is around 1% to 2% while the ZIO Patch is 66%.
- The ZIO Patch is best used in individuals with a history of frequently undiagnosed syncope when episodes are likely to occur during the 14 day monitoring period.
- Implantable loop recorders are suggested as a first-line diagnostic tool for patients lacking classical etiological features.

Keywords: Arrhythmia Detection with Long Term Continuous ECG Monitoring

Retrospective data from 26,751 consecutive patients undergoing first-time ZIO Patch studies during 2011 were analyzed for wear time, analyzable signal time, diagnostic yield and timing of arrhythmia detection.

- The mean wear time was 8 days and the median analyzable time was 99% of the total wear time.
- After 48 hours, the ZIO Service detected:
  - 51% of patients had their first symptom-triggered arrhythmia
  - 47% of patients experienced their first symptomatic episode of atrial fibrillation
  - 37% of patients had their first symptomatic episode with AV block
  - 30% of patients had their first arrhythmia of any type


Keywords: ZIO Patch Compared to Holter, Change in Clinical Management for AFib

Prospective 74-patient study in which each patient simultaneously wore a Holter Monitor for 24 hours and the ZIO Patch for an average of 10.8 days.

- Over a 24-hour period, the ZIO Patch and Holter monitor equally identified AFib events and estimated AFib burden.
- However, following ZIO monitoring, AFib events were identified in 18 additional individuals and the documented pattern of AFib changed in 21 patients. Additionally, potentially malignant arrhythmias were first recorded on the ZIO Patch after 24 hours of monitoring.
- Longer continuous monitoring with ZIO resulted in a meaningful change in clinical management for 28.4% of patients.