

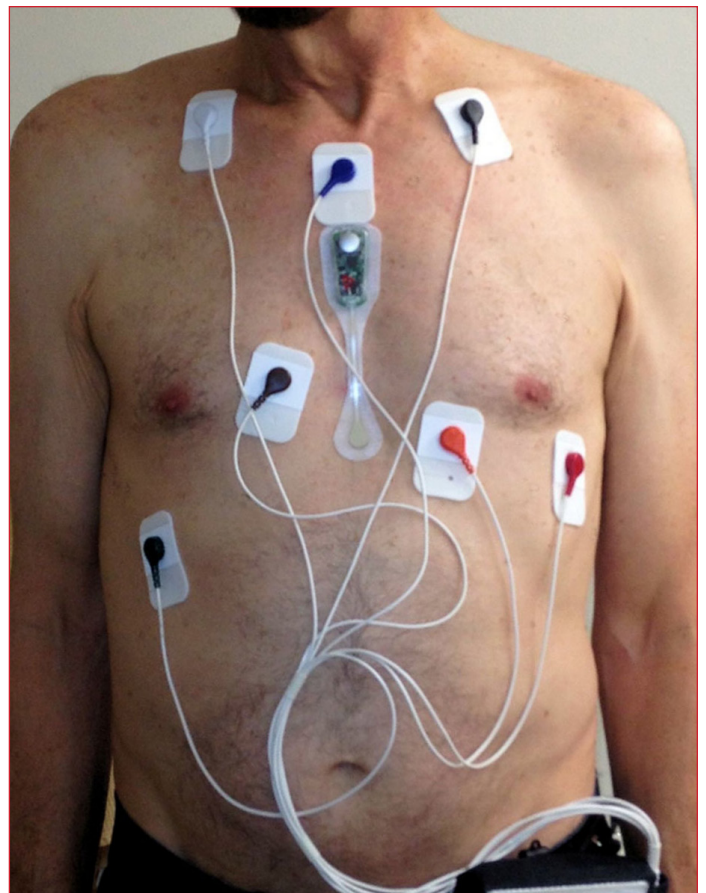
CAM vs. Holter Study

STUDY PURPOSE

To compare simultaneous recordings to determine diagnostic efficacy between an external patch system specifically designed to ensure better P-wave recordings and a standard Holter monitor

STUDY METHODS

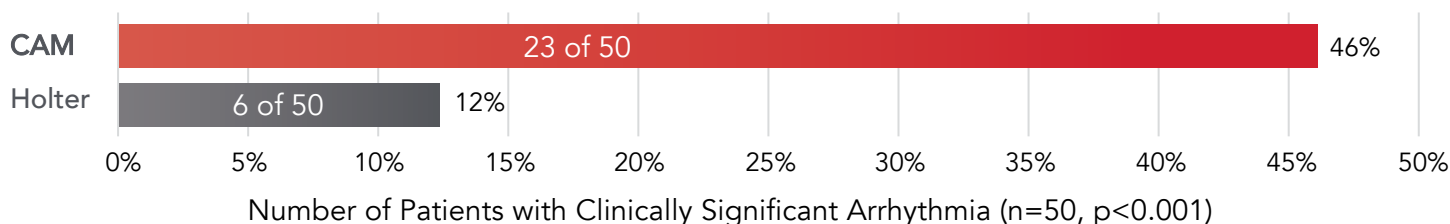
- Prospective comparison of a single-channel patch monitor and a standard 3-lead Holter monitor:
 - **Carnation Ambulatory Monitor (CAM)** (Bardy Diagnostics, Inc.)
 - Standard DR180 Series 3-channel (leads V1, II, and V5) Holter monitor (NorthEast Monitoring, Inc.)
- 50 consecutive adult patients enrolled from a single center:
 - Both devices simultaneously applied and removed after 24 hours
 - Each patient served as their own control
- Holter and **CAM** reports were read in a blinded fashion by two electrophysiologists unaware of the findings in the other corresponding ECG recording
- All patients, technicians, and physicians completed a questionnaire on comfort, ease-of-use, and potential complications



	OUTCOME MEASURES
Primary	Impact on Clinical Decision-Making When Comparing Rhythm Findings
Secondary	- Patient Assessment - Device Preference - Comfort - Skin Irritation - Discreetness - Effect on Daily Activities - Effect on Sleeping - Clinician Assessment - Device Stability - Ease of Attachment

STUDY RESULTS

IMPACT ON CLINICAL DECISION-MAKING } The **CAM** Patch yielded clinically significant information that either altered patient management and/or prevented the need for intervention as indicated by the Holter.



The **CAM** Patch identified arrhythmias missed or misidentified by the Holter in over a third of the patients. The Holter identified only a subset of clinically significant arrhythmias, all of which were also found using the **CAM** Patch.

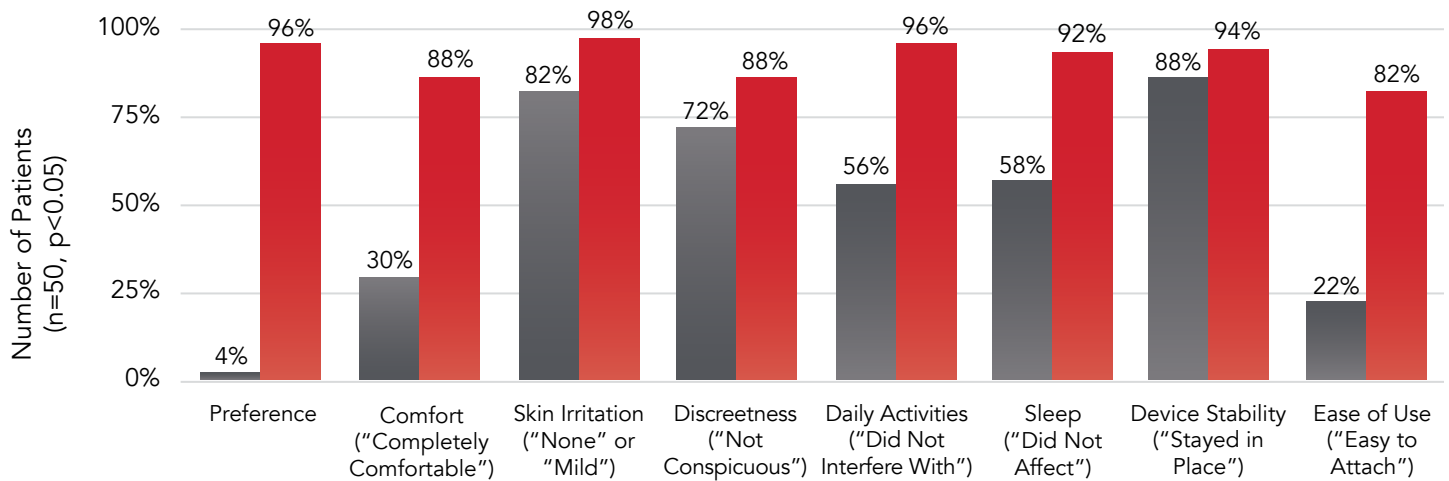
Missed by Holter (7 of 50 patients)	Misidentified by Holter (10 of 50 patients)	Identified by Both (6 of 50 patients)
- AFI, in addition to AF – Identified as AF only on Holter (3 patients) - NSVT - Sinus Arrest - AVB - CHB and Sinus Arrest*	- AT – Identified as AF on Holter (2 patients) - ST – Identified as AT on Holter - No AF – Identified as AF on Holter - AF – Identified as AT on Holter - ST – Identified as AT on Holter - No PVCs – Identified noise as frequent PVCs on Holter 1:1 AT – Identified as ST on Holter - AT with no VT – Identified as AF with VT on Holter - AFI with CHB – Identified as AF with junctional escapes on Holter	- NSVT (2 patients) - Sinus Arrest - AF & AFI - AF - Wenckebach AVB

* 1 pt had 2 arrhythmias missed

Abbreviations: AF, atrial fibrillation; AFI, atrial flutter; AT, atrial tachycardia; AVB, atrioventricular block; CHB, complete heart block; NSVT, nonsustained ventricular tachycardia; PVCs, premature ventricular contractions; ST, sinus tachycardia.

PATIENT & CLINICIAN ASSESSMENT

The **CAM** Patch outperformed the Holter monitor on all comparison metrics. The **CAM** Patch was significantly preferred by patients over the Holter monitor.



STUDY CONCLUSION



In a direct comparison on 50 consecutive patients, the single-channel **CAM** Patch monitor demonstrated to be comfortable, easy-to-use, and designed to reliably capture the P-wave as compared to the Holter monitor. As a result of the superior ECG clarity, it resulted in significantly improved rhythm diagnoses when compared to the standard 3-lead Holter.

The Carnation Ambulatory Monitor is designed to provide extended-duration cardiac monitoring for people who are suspected of having cardiac arrhythmias. **Rx only.** For safe and proper use of the products mentioned herein, please refer to the Instructions for Use.

Source: Smith WM, Riddell F, Madon M, Gleva MJ. Comparison of diagnostic value using a small, single channel, P-wave centric sternal ECG monitoring patch with a standard 3-lead Holter system over 24 hours. American Heart Journal. 2017;185:67-73. doi:10.1016/j.ahj.2016.11.006

BardyDx.com | Baxter.com

Baxter, BardyDx, BDx design, Bardy Diagnostics and CAM are trademarks of Baxter International Inc. or its subsidiaries. Any other trademarks, product names or brand images appearing herein are the property of their respective owners.

US-FLC199-220019 (v7.0) 05/2025