

Detection of Atrial Fibrillation Recurrence and Monitoring Duration on Ambulatory Cardiac Monitoring

Implications for OCEAN Trial-Guided Anticoagulation Discontinuation

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BACKGROUND

- Atrial fibrillation (AF) recurrence is common after ablation.
- OCEAN Trial showed anticoagulation may be stopped if no AF detected post-ablation, based on sequential Holter monitoring of 24-48 hours.
- Short-duration monitoring may miss recurrence.

METHODS

- US cohort (2018-2022) receiving Zio ambulatory cardiac monitor (ACM)[†] and with AF ablation (CPT/ICD-10 PCS code-based) within the prior year
- AF subtype (ICD-10 code-based): paroxysmal (PAF), persistent/permanent (PerAF), unspecified
- ECG data analyzed using an FDA-cleared deep learning algorithm and confirmed by qualified cardiographic technicians
- Tokenized linkage to claims for commercial fee-for-service and government sponsored plans, including Medicare Advantage

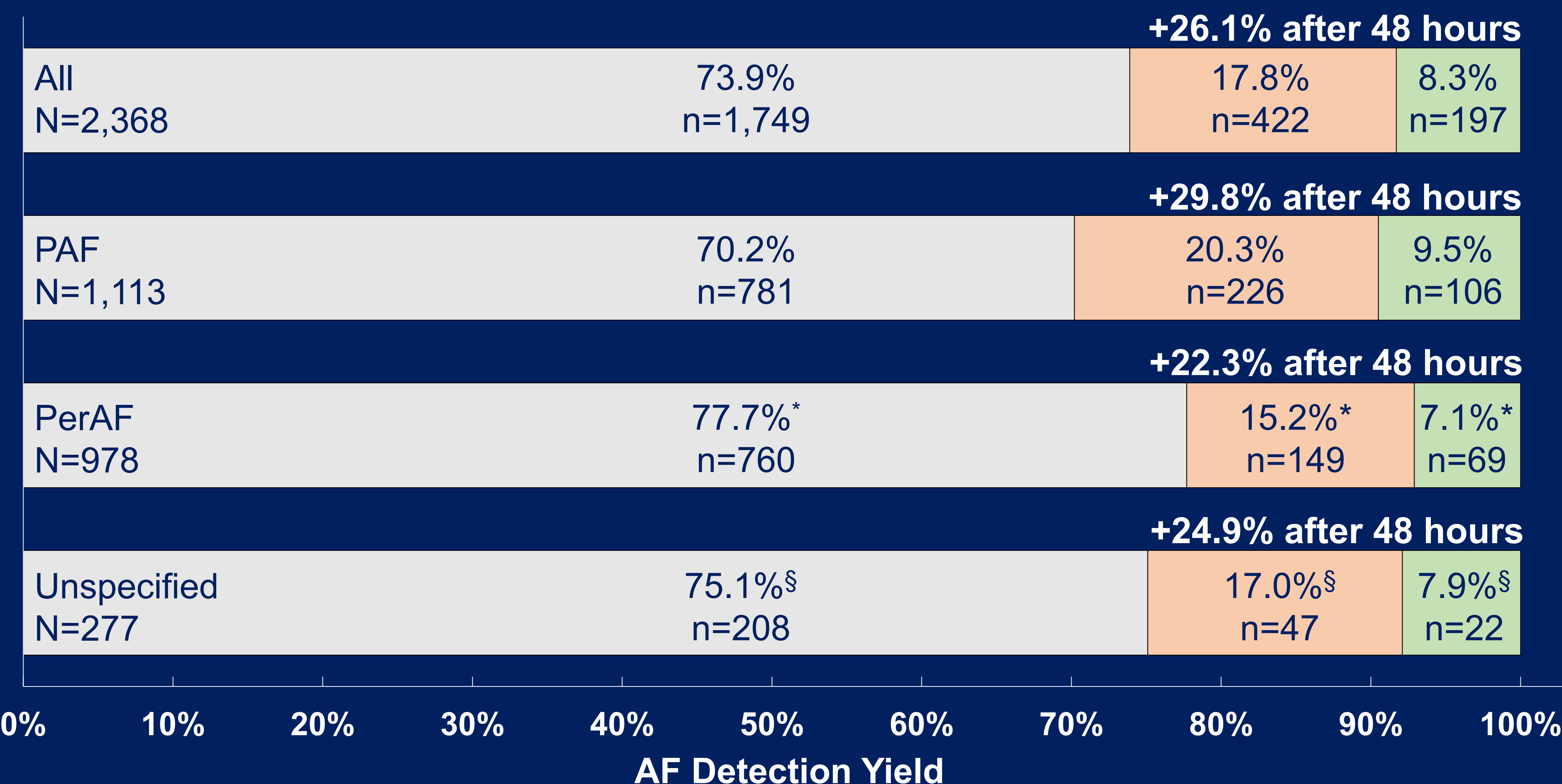
RESULTS

- 709,083 patients with linked data
- N=11,051 met inclusion criteria
- 53% PAF, 36% PerAF, 11% Unspecified AF
- Age: Mean (SD) Age 65.6 (10.2) years
- 39% female
- 74% White race; 26% Black/Asian/Other/Unknown
- CHA₂DS₂-VASc Stroke Risk: Mean (SD) 2.4 (1.5) Overall
- Median time to monitoring: 113 days post-ablation

Monitoring for ≤48 hours misses ~25-30% of AF recurrence.

Timeline of Post-Ablation AF Recurrence

■ 0-2 days ■ >2-7 days ■ >7-14 days



*p<0.05 vs PAF §p=NS vs PAF

[†]Adults ≥18 years undergoing ≤14 days of continuous patch-based ACM (Zio[®] XT or AT; iRhythm Technologies, San Francisco, CA) and with AF ablation (CPT/ICD-10 PCS code-based) within the prior year.

Missed AF may lead to misclassification of patients being considered for anticoagulation discontinuation under an OCEAN-guided approach.

AF RECURRENCE, BURDEN, AND REPEAT ABLATION

| | All (N=11,051) | PAF (n=5,812) | PerAF (n=4,012) | Unspecified (n=1,227) |
|------------------------------|-------------------|------------------|--------------------|--------------------------|
| ACM detected AF ≥ 30s | | | | |
| Overall (n, % of AF Group) | 2,368 (21.4%) | 1,113 (19.2%) | 978 (24.4%)* | 277(22.6%)* |
| AF Burden[†] | | | | |
| Median (IQR) | 11.9% (87.8%) | 4.4% (22.1%) | 43.1% (94.9%)* | 16.8% (97.9%)* |
| Repeat Ablations | | | | |
| 2018-2021 ablations (n) | 8,380 | 4,429 | 3,024 | 927 |
| Repeat ablation within 1 yr | 667 (8.0%) | 318 (7.2%) | 265 (8.8%)* | 84 (9.1%)* |

*p<0.05 vs PAF [†]Among patients with non-zero burden

KEY FINDINGS

- 21.4% of patients had AF recurrence during monitoring.
- 8.0% received a repeat AF ablation within one year.
- When compared to 14 days of continuous, uninterrupted ACM, an OCEAN-guided approach of up to 48 hours of monitoring would miss 29.8% of AF in paroxysmal and 22.3% of AF in persistent patients.
- Among patients with recurrence, AF burden was substantial and varied by subtype.

CLINICAL IMPLICATIONS

- Monitoring duration directly impacts anticoagulation decision-making.
- Short-duration (≤48 hours) monitoring may miss clinically meaningful AF recurrence.
- Extended (e.g., 14 days continuous, uninterrupted) monitoring may improve patient classification post-ablation.

LIMITATIONS

- Retrospective observational design
- AF types were identified using administrative claims, which may result in misclassification
- Variability in timing of post-ablation monitoring

DISCLOSURES

- Dr. Turakhia has received equity from iRhythm, Connect America, Evidently, PocketRN, AliveCor, and Hippocratic.ai. Dr Turakhia is an employee and corporate officer of iRhythm Technologies Inc.
- S Schmitt has received compensation from iRhythm.
- V Fokin, JM Ashburner, and AJ Battisti are employees of and have received equity from iRhythm Technologies, Inc.