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Data in Brief





Data Article

Initial experience with the convergent procedure for longstanding persistent atrial fibrillation: A 5 year dataset



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ABSTRACT

In patients with longstanding persistent atrial fibrillation (AF), outcomes from catheter ablation remain suboptimal. The convergent procedure combines minimally invasive surgical ablation with subsequent catheter ablation, and may contribute towards maintenance of sinus rhythm in this patient group. We performed the convergent procedure on 43 patients with longstanding persistent AF from 2013-2018. Patients underwent clinical review at 3, 6, and 12 months and thereafter as necessitated by their symptoms. Our dataset describes patients' baseline characteristics and rhythm control protocols, as well as outcomes including arrhythmia recurrence, the need for antiarrhythmic drugs, requirement for repeat rhythm control procedures, and complications. These data provide a real world insight into the risks and benefits of the convergent procedure in patients with longstanding persistent AF.

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Specifications table

Subject Cardiology and Cardiovascular Medicine Specific subject area Electrophysiology: the convergent procedure (hybrid surgical ablation) to treat longstanding persistent AF Type of data Tables of raw data Kaplan-Meier plot How data were acquired How data were acquired	
longstanding persistent AF Type of data Tables of raw data Kaplan-Meier plot	
Type of data Tables of raw data Kaplan-Meier plot	
Kaplan-Meier plot	
records	
Data format Summated outcome data in Table 1	
	Survival distribution (Kaplan-Meier plot) in Fig. 1
Raw numerical and categorical baseline and follow-up data in supplementary Table S1	
(see supplementary file)	
Parameters for data Baseline parameters: age, gender, hypertension (HTN), diabetes, ischaemic heart	
collection disease (previous percutaneous coronary intervention), current or previous smoking	
history, echocardiographic data, duration of atrial fibrillation, anti-arrhythmic drug	
use, anticoagulation strategy, previous ablation or cardioversion (DCCV), pacemaker,	
history of hypertrophic cardiomyopathy, symptomatic class according to EHRA and	
NYHA guidelines	
Catheter ablation data: number of ablations required, ablation lines delivered,	
complications	
Surgical ablation data: access used, number of ablation lesions delivered, complications	
Follow-up data: repeat echocardiographic data, repeat symptomatic data, rhythm data	
according to clinic electrocardiograms (ECGs), holter monitor examination and/or	
pacemaker interrogation	
Description of data Table 1:	
collection - Summated outcomes for all patients according to procedure reports and clinical	
records	
Fig. 1:	
- Kaplan-Meier plot showing arrhythmia-free survival over time for patients	
undergoing the convergent procedure (allowing for multiple catheter ablations and	
anti-arrhythmic drugs) Supplementary Table S1:	
- Procedural data were extracted from operation notes and procedure reports	
- Rhythm data were obtained from 72 h holter monitors, clinic ECGs and pacemaker	
interrogations where available	
- Echocardiographic data were extracted from the hospital's imaging database	
- Symptomatic data and medical co-morbidities were discerned from clinic visits and	
electronic medical records	
Data source location City/Town/Region: St Bartholomew's Hospital, London	
Country: UK	
Data accessibility With the article:	
- Table 1: within the article	
- Fig. 1: within the article	
- Table S1: supplementary file	
Related research article E Maclean, J Yap, B Saberwal, S Kolvekar, W Lim, N Wijesuriya, N Papageorgiou, G	
Dhillon, RJ Hunter, M Lowe, P Lambiase, A Chow, H Abbas, R Schilling, E Rowland, S	
Ahsan	
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atrial fibrillation: a single centre, propensity-matched cohort study	
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Value of the data

• These data evaluate an emerging hybrid ablation technique for treating atrial fibrillation (AF). No randomised controlled trial data has yet been published on this subject, and previous observational data demonstrate significant heterogeneity regarding procedural success and the incidence of complications [2,3]. Our results include outcomes for individuals with very longstanding persistent AF; there is currently only limited published data for this patient group.

Table 1Clinical outcomes (at 1 year and long-term) for patients undergoing the convergent procedure.

Outcome	Patients undergoing the convergent procedure $(n = 43)$	
AF-free at 1 year (single procedure, on AADs)	60.5% (n = 26)	
AF-free at 1 year (single procedure, off AADs)	37.2% (n = 16)	
Arrhythmia-free long-term (multiple procedures, on AADs; mean follow-up 30.5±13.3 months)	58.1% (n = 25)	
Arrhythmia-free long-term (multiple procedures, off AADs; mean follow-up 30.5±13.3 months)	32.5% (n = 14)	
Off AADs at last follow-up	60.5% (n = 26)	
New atrial tachycardia	32.6% n = 14)	
Complications	11.6% (n = 5)	
	Tamponade (pericardiocentesis)	2
	Emergency sternotomy	1
	Pericardial hernia	1
	Phrenic nerve palsy	1

- The convergent procedure requires ongoing validation as a potential rhythm control strategy. Our data examines procedural success both at 1 year post procedure and long term, and includes a breakdown of complications. A full description of the risks and the efficacy of the procedure can inform clinicians as to its suitability as a treatment option for their patients.
- Our observational data can be incorporated into other registries to potentiate more largescale analyses of the utility of the convergent procedure. By publishing our raw data in its entirety on a per-patient basis, researchers also have access to specific data subgroups, such as those individuals with pacemakers, cardiomyopathy or severe systolic dysfunction.

1. Data

The dataset contains raw numerical and categorical data for all patients at the point of recruitment and during follow-up. This includes baseline characteristics such as age, gender and duration of AF, echocardiographic and symptomatic data, and history of previous rhythm control procedures. The presenting rhythm is documented for each follow-up visit, and the total follow-up duration listed in months. Anti-arrhythmic drug use is described according to class of action. The latest echocardiographic and symptomatic data is also listed, and procedural complications are itemised. These data are displayed in supplementary Table S1.

Outcome data is summarised according to the primary and secondary outcomes, and is subdivided by duration of follow-up and the requirement for anti-arrhythmic drugs. Incidence of atrial tachycardia is also documented. These findings are described in Table 1, with long-term arrhythmia free survival visualised in Fig. 1.

2. Experimental design, materials, and methods

2.1. Ethics

This project was registered with the local clinical effectiveness unit. As a retrospective analysis of patient records, the need for formal ethical approval was waived.

2.2. Methods

Our method for performing the convergent procedure has been described in detail elsewhere [1]. 43 patients underwent the procedure via a staged approach. Following the second part of

Arrhythmia-free survival for patients undergoing the Convergent Procedure (allowing for multiple catheter ablations and AADs)

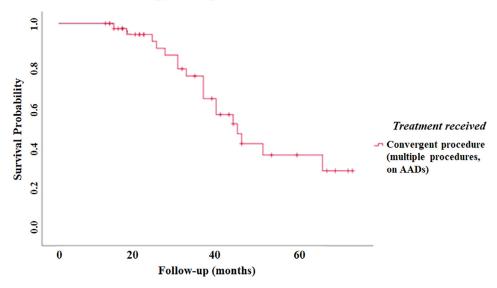


Fig. 1. Kaplan-Meier plot showing arrhythmia-free survival over time for patients undergoing the convergent procedure (allowing for multiple catheter ablations and AADs).

the procedure (catheter ablation), a 3 month blanking period was observed in line with international consensus recommendations [4]. Subsequent clinical review took place at 3 months (with ECG), 6 months (with 72 h holter analysis) and 12 months (with echocardiogram, ECG and symptom-guided 72 h holter monitor). Patients with pacemakers in situ underwent device interrogation in addition to holter monitoring. Further clinical follow-up took place annually or sooner if warranted by symptoms.

2.3. Outcomes

A recurrence was defined as more than 30 s of documented AF outside of the blanking period. The presence of any other atrial arrhythmia was also recorded. Anti-arrhythmic drug (AAD) use was assessed at 12 months, and anticoagulation was continued as indicated by CHA₂DS₂VASc score. The primary outcome was AF-free survival at 12 months. Secondary outcomes included incidence of atrial tachycardia, change in NYHA and EHRA class, procedural complications, echocardiographic data, freedom from AADs, and arrhythmia-free survival long term. Patients underwent additional DCCV or repeat catheter ablation as indicated; in these cases, a further 3 month blanking period was observed and follow-up restarted.

Conflict of Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.dib.2020.105417.

References

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