# Impact of contact force technology on reducing the recurrence and major complications of atrial fibrillation ablation: A systematic review and meta-analysis

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## Abstract

Contact force (CF) monitoring can be useful in accomplishing circumferential pulmonary vein (PV) isolation for atrial fibrillation (AF). This metaanalysis aimed to assess the efficacy and safety of a CF-sensing catheter in treating AF. Randomized controlled trials or non-randomized observational studies comparing AF ablation using CF-sensing or standard non-CF (NCF)-sensing catheters were identified from PubMed, EMBASE, Cochrane Library, Wanfang Data, and China National Knowledge Infrastructure (January 1, 1998–2016). A total of 19 studies were included. The primary efficacy endpoint was AF recurrence within 12 months, which significantly improved using CF-sensing catheters compared with using NCF-sensing catheters [31.1% vs. 40.5%; risk ratio (RR)=0.82; 95% confidence interval (CI), 0.73–0.93; p<0.05]. Further, the acute PV reconnection (10.1% vs. 24.2%; RR=0.45; 95% CI, 0.32–0.63; p<0.05) and incidence of major complications (1.8% vs. 3.1%; OR=0.59; 95% CI, 0.37–0.95; p<0.05) significantly improved using CF-sensing catheters compared with using NCF-sensing catheters. Procedure parameters such as procedure duration [mean difference (MD)=–28.35; 95% CI, -39.54 to –17.16; p<0.05], ablation time (MD=–3.8; 95% CI, -6.6 to –1.0; p<0.05), fluoroscopy duration (MD=–8.18; 95% CI, -14.11 to -2.24; p<0.05), and radiation dose (standard MD=–0.75; 95% CI, -1.32 to -0.18; p<0.05] significantly reduced using CF-sensing catheters. CF-sensing catheter ablation of AF can reduce the incidence of major complications and generate better outcomes compared with NCF-sensing catheters during the 12-month follow-up period. (*Anatol J Cardiol 2017; 17: 82-91*) **Keywords:** atrial fibrillation; ablation; contact force-sensing catheter; meta-analysis

## Introduction

Atrial fibrillation (AF) is the most common sustained arrhythmia, and ablation procedures for AF have been shown to be safe and effective in a large number of cases worldwide (1–4). However, the recurrence rates of AF after catheter ablation are still considerably high (5, 6). Pulmonary vein (PV) reconnection due to ineffective ablation lesions has been identified as the main cause of AF recurrence (7, 8), and catheter–tissue contact is essential for effective ablation lesions (4, 9, 10). However, an accurate measurement of lesions and understanding the limitations of the contact force (CF) are crucial for avoiding complications (11). In recent years, radiofrequency (RF) catheter ablation with CF sensing, a novel method, has been claimed to be potentially responsible for effective ablation. When using it, the catheter–tissue CF can be measured at the catheter tip with fiber optic or magnetic sensors (12).

The safety and effectiveness of CF-sensing catheters have been evaluated in ex vivo models (10, 13) and in vivo experimental studies (14, 15) before their recent application in humans. Experimental data in previous studies have demonstrated a strong relationship between CF and lesion size when using an RF current for catheter ablation (14). However, the efficacy and safety of CF-sensing catheters, particularly for reducing the rate of complications, remain controversial.

The purpose of this meta-analysis was to evaluate the efficacy and safety of catheter AF ablation using CF-sensing catheters.

## Methods

## Literature search

Electronic databases, such as PubMed, EMBASE, Wanfang Data, China National Knowledge Infrastructure (January 1, 1998–2016), and Cochrane Controlled Trials Register, for reports on all randomized controlled trials (RCTs) or non-randomized observational studies (NROSs) published in English or Chinese were searched using the following medical subject headings, "contact force-sensing catheter," "ablation," and "atrial fibrillation," to capture data on catheter AF ablation using CF-sensing



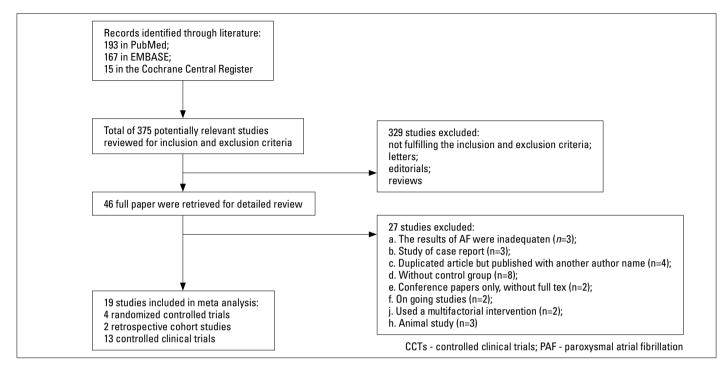


Figure 1. Flow diagram of the literature search stages

catheters. The abstracts of all identified RCTs or NROSs were independently screened by two reviewers.

## Study selection and quality assessment

Studies fulfilling the following criteria were included: (1) patients undergoing AF ablation using CF-sensing catheters and standard non-CF (NCF)-sensing catheters, (2) patients with paroxysmal AF (PAF) or persistent AF (Per AF), and (3) human studies conducted in adults who were 18 years and older. Non-comparative trials, case reports, editorials, and reviews were excluded from this study.

We used PRISMA guidelines in this meta-analysis. Individual studies were checked for the following characteristics: adequate sequence generation, allocation concealment, attrition less than 15%, blinded assessment, intent-to-treat analysis, complete follow-up, and adequate AF monitoring.

## **Data abstraction**

The citations were also reviewed, and data were independently abstracted by two reviewers; disagreements were resolved by discussions. Abstracted data included the following: (1) study type, study size, study design, CF catheter used, mean CF used, and follow-up; (2) age and gender; (3) AF recurrence within 12 months (primary outcomes); (4) occurrence of acute PV reconnection; (5) primary safety endpoint including device-related serious adverse events (events were classified as major and minor complications; major complications included in-hospital death, cardiac perforation, cardiac effusion or tamponade, stroke, PV stenosis, esophageal fistula, severe hemoptysis, phrenic nerve lesion, and thromboembolic event, whereas minor complications were mainly related to vascular access complications, including femoral/subclavian hematoma and arteriovenous fistula); and (6) procedure duration, ablation time, fluoroscopy duration, and radiation dose.

#### **Statistical analysis**

Statistical analysis was performed using Cochrane RevMan version 5 (The Cochrane Collaboration, UK), and results were expressed as weighted mean differences (MDs) and relative risk for continuous and dichotomous outcomes, respectively, with 95% confidence intervals (CIs). Outcomes were pooled using the random-effects model when the heterogeneity was moderate or high ( $l^2$ >50%). However, the fixed-effects model was used when the heterogeneity was low ( $l^2$ <50%). Radiation doses used among the included studies were compared using a standard MD (SMD) as different radiation units had been used. The present study assessed the heterogeneity between studies using the Cochran's Q statistic and  $l^2$  index. All statistical testing was two tailed with statistical significance at p<0.05.

#### Results

The electronic search identified 193 references from PubMed, 167 from EMBASE, and 15 from the Cochrane Central Register of Controlled Trials. Among these abstracts, 329 were excluded. The full manuscripts for the remaining 46 studies were retrieved for a detailed review, and 27 were further excluded. Finally, 19 studies (16–34) [4 RCTs (16–19), 2 retrospective cohort studies (20, 21), and 13 NROSs (22–34)] were identified that compared the safety and efficacy of CF-sensing or NCF-sensing catheters in the setting of AF ablation. Information relevant to the literature search is shown in Figure 1.

## **Publication bias**

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No significant publication bias was found for the primary outcome (AF recurrence at the follow-up) as assessed by a funnel plot (Fig. 2).

#### **Baseline patient characteristics**

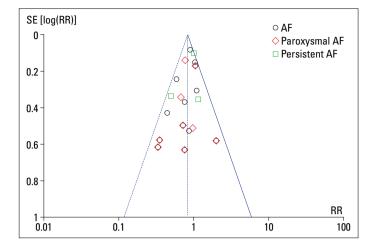
Baseline patient characteristics are provided in Table 1. A total of 4053 patients were included in the CF-sensing (n=1546) and NCF-sensing (n=2507) catheter groups.

Ten studies provided detailed information on the PAF and/ or Per AF patient subgroups, and relevant information was abstracted to compare the efficacy and safety in the AF, PAF, and/or Per AF subgroups.

## Efficacy of AF ablation using CF-sensing catheters

AF recurrence within 12 months was compared in the AF (13 studies), PAF (9 studies), and Per AF (3 studies) subgroups. In

Contact force-guided ablation Standard radiofrequency ablation



**Figure 2.** Funnel plot for the assessment of publication bias for the primary outcome. Effect size is plotted on the x -axis and SE on the y-axis. AF - atrial fibrillation; RR - risk ratio; SE - standard error

**Risk ratio** 

Study or subgroup Events To	tal Events Tota	al Weight M–H, Fixed, 95% C	I Year M–H, Fixed, 95% Cl
Jarman 2014         100         2           Wutzler 2014         5         5           Marijon 2014         3         5           Sciarra 2014         5         1           Ullah 2014         32         13           Wakili 2014         13         1           Itoh 2015         2         Nakamura 2015         6           Sigmund 2015         9         9         8           Reddy 2015         49         1         1	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2014
Itoh 2015 2	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	50 1.8% 0.35 [0.11, 1.09] 21 0.9% 0.97 [0.36, 2.66] 54 2.6% 0.67 [0.34, 1.31] 50 1.4% 0.22 [0.05, 0.98] 13 7.2% 1.05 [0.75, 1.47]	2014 2014 2014 2014 2014 2014 2014 2015 2015 2015 •
Jarman 2014 62 1 Sigmund 2015 9	08 124 21 37 17 3 <b>59 26</b> 148	14 1.1% 1.14 [0.57.2.29] 16 13.1% 1.00 [0.82, 1.22] 25 2.8% 0.50 [0.26, 0.97] 25 <b>17.0% 0.93 [0.77, 1.12]</b>	2014 2014 2015
Total (95% Cl)         14           Tatol events         448           Heterogeneity: Chi²=34.52, df=24 (P=0.08) l²=30°         12=30°           Test for overall effect: Z=4.28 (P<0.0001)	817	33 100.0% 0.82 [0.75, 0.90]	0.01 0.1 1 10 100 Favours [Contact force-guided ablation] Favours [Standard radiofrequency ablation]
<b>b</b> Contact force-quided ablat	ion Standard radiofrequency abla	ation Risk ratio	Risk ratio
Study or subgroup Events To	tal Events Tota	al Weight M–H. Fixed, 95% Cl	Year M–H, Fixed, 95% Cl
Reddy 2015 16 1 Nakamura 2015 7	83 9 8 30 5 3 25 26 5 52 20 14 60 16 6	60 17.6% 0.44 [0.19, 0.99]	2012
Total (95% CI)         3           Tatol events         40           Heterogeneity: Chi?=5.55, df=6 (P=0.48); l²=0%           Test for overall effect: Z=4.59 (P<0.00001)	<b>95 40</b> 99	<b>)9 100.0% 0.45 [0.32, 0.63]</b>	0.01 0.1 1 10 100 Favours [Contact force-guided ablation]

**Risk ratio** 

Figure 3. (a) Forest plot showing the RR and 95% CI for AF recurrence within 12 months for studies comparing the CF and NCF groups. (b) Forest plot showing the RR and 95% CI for the occurrence of acute PV reconnection for studies comparing the CF and NCF groups

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Follow up months	13	12	NR	12	12	12	NR	9	12	12	2.5	12	13.2±0.9	12
Mean CF, g	NR	18	17.8	NR	16	NR	NR	NR	16	13	NR	17.4	NR	26.8
CF Catheter	TactiCath	Thermocool SmartTouch	Thermocool SmartTouch	Thermocool SmartTouch	Thermocool SmartTouch	Thermocool SmartTouch	Thermocool SmartTouch	Thermocool SmartTouch	Tacticath	Thermocool SmartTouch	Thermocool SmartTouch	Tacticath	Thermocool SmartTouch	TactiCath
EF (%) (CF/NCF)	62.4±7.1 /62.4±6.2	67/65	56.0±7.9 /58.1±8.0	65±10 /65±7	60±7 /60±6	56±5 /57±7	NR	65.7±5.2 /62.4±11.8	62.3±7.4 /62.0±7.8	NR	56±5 ∕55±5	68.5±2.2 /65.0±1.9	63.3±5.5 ∕59.9±5.4	56.8±4.9 /55.6±3.1
LA size mm (CF/NCF)	39.9±5.9 /39.3±4.5	40±6/39±5	42.0±3.6 /43.0±4.3	37±7 /38±6	44±6 /45±6	40±6 ∕41±6	NR	41.3±7.8 /42.0±6.8	43.2±6.4 /41.9±5.5	4.4±0.6 /4.4±0.6	35±7 /36±6	43.2±0.9 /42.1±0.9	32.4±14.2 /39.2±4.7	41.5±6.1 /42.4±6.7
Diabetes, n(%) (CF/NCF)	16 (10.5) /17 (11.9)	8 (13.3) /10 (16.7)	2 (8.3) /0 (0)	5 (10) /8 (16)	4(11) /4 (11)	4 (4) /3 (3)	31 (6) /50 (5)	3 (15.8) /4 (21.1)	RN	3 (5) /2 (4)	1 (5) /2 (10)	NR	NR	3 (9.7) /10 (8.9)
Hypertension n(%) (CF/NCF)	75 (49.3) /69 (48.3)	27 (45.0) /36 (60.0)	8 (33.3) /6 (50.0)	32 (64) /26 (52)	25 (71) /29 (83)	46 (47) /52 (53)	77 (15) /140 (14)	13 (68.4) /9 (47.4)	6 (30) /12 (34)	11 (22) /7 (14)	NR	21 (65.6) /25 (71.4)	NR	20 (64.5) /58 (51.8)
Male, n(%) (CF/NCF)	100 (65.8) /91 (63.6)	44 (73.3) /45 (75.0)	19 (79.2) /11 (91.7)	30 (60) /31 (62)	24 (69) /27 (77)	71 (72) /68 (69)	349 (68.4) /264 (63.6)	12 (63) /17 (89)	16 (80) /28 (80)	41 (82) /39 (78)	18 (86) /18 (86)	21 (65.6) /23 (65.7)	19 (76) /43 (86)	21 (67.7) /71 (63.4)
Mean age y(CF/NCF)	59.6±9.3 /61.0±10.8	64/64.5	58.6±11.3 /62.2±8.5	65±11 /61±10	67±9 /60±11	59.5±9.6 /59.5±9.4	60.5±11.0 /60.8±11.3	62.5±10.1 /57.3±8.6	58±10 /56±13	63/62	59.7±9.1 /54.6±11.0	63.6±1.7 /59.3±1.9	58.8±12.7 /58.6±11.0	59.8±10.9 /60.9±10.2
PerAF (CF/NCF)	0	40 (22/18)	9 (6/3)	0	26 (16/10)	72 (37/35)	750 (255/495)	10 (4/6)	0	NR	0	28 (14/14)	0	39 (12/27)
PAF (CF/NCF)	295 (152/143)	80 (38/42)	27 (18/9)	100 (50/50)	44 (19/25)	126 (62/64)	656 (238/418)	28 (15/13)	55 (20/35)	NR	42 (21/21)	39 (18/21)	75 (25/50)	104 (19/85)
AF (CF/NCF)	295 (152/143)	120 (60/60)	36 (24/12)	100 (50/50)	70 (35/35)	198 (99/99)	1515 (510/1005)	38 (19/19)	55 (20/35)	100 (50/50)	42 (21/21)	67 (32/35)	75 (25/50)	143 (31/112)
Type of study	prospective, randomized, controlled, multicenter study	prospective, randomized, controlled study	Prospective non-randomized study	Prospective non-randomized study	Prospective non-randomized study	Prospective case-matched control trial	retrospective observational cohort study	prospective, randomized, controlled, study	prospective, randomized, controlled, study	Prospective non-randomized multicenter study	Prospective non-randomized study	Prospective non-randomized study	Prospective non-randomized study	Prospective non-randomized study
	Reddy 2015 (TOCCASTAR)	Nakamura 2015	Wolf 2015	ltoh 2015	Makimoto 2015	Sigmund 2015	G. Lee 2015	Kimura 2014	Casella 2014	Ullah 2014	Sciarra 2014	Wakili 2014	Andrade 2014	Wutzler 2014

Table 1. Baseline clinical characteristics and follow-up of the patients

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	Type of study	AF (CF/NCF)	PAF (CF/NCF)	PerAF (CF/NCF)	Mean age y(CF/NCF)	Male, n(%) (CF/NCF)	Hypertension n(%) (CF/NCF)	Diabetes, n(%) (CF/NCF)	LA size mm (CF/NCF)	EF (%) (CF/NCF)	CF Catheter	Mean CF, g	Follow up months
Marijon 2014	Prospective non-randomized study	60 (30/30)	60 (30/30)	0	59.9±9 /61.0±10	21 (70.0) /22 (73.3)	NR	NR	NR	64.7±4 /65.4±5	Thermocool SmartTouch	21.7	12
Akca 2014	Prospective non-randomized study	449 (143/306)	NR	NR	55.7±15.1 /51.7±16.6	NR	NR	NR	NR	NR	Thermocool SmartTouch and Tacticath	NR	NR
Jarman 2014	Retrospective case- control study	600 (200/400)	276 (92/184)	324 (108/216)	63±12 /61±10	149(74.5) /282(70.5)	80 (40) /119 (30)	21 (11) /34 (9)	42±7 /44±7	NR	Thermocool SmartTouch	NR	<b>11.4±4.7</b>
Haldar 2012	Prospective non-randomized study	40 (20/20)	14 (7/7)	26 (13/13)	63±14 /61±12	15 (75) /11 (55)	7 (35) /6 (30)	NR	42±8 ∕41±5	57±12 /59±10	Thermocool SmartTouch	NR	NR
Martinek 2012	Prospective non-randomized study	50 (25/25)	50 (25/25)	0	60.5±9.5 /57.4±11.6	12 (48) /17 (68)	10 (40) /12 (48)	3 (12) /1 (4)	39±6 /37±6	53±4 /53±3	Thermocool SmartTouch	NR	NR
Mean												18.3	
Total		4053 (1546/2507)	2071 (849/1222)	1324 (487/837)									
AF - atrial fibrillation	AF - atrial fibrillation; CF - contact force; NR - not reported; PAF - paroxysmal atrial	not reported; PAF	: - paroxysmal a		Per AF - persist	ent atrial fibrillati	on. Statistical analys	fibrillation; Per AF - persistent atrial fibrillation. Statistical analysis was performed using the Cochrane RevMan version 5 software	ng the Cochrane Rev	Man version 5 s	software		

the AF and PAF subgroups, AF recurrence significantly improved using CF-sensing catheters compared with that using NCF-sensing catheters in the AF [31.1% vs. 40.5%; risk ratio (RR)=0.82; 95% CI, 0.73–0.93; I<sup>2</sup>=32%; p=0.001] and PAF (25.3% vs. 40.0%; RR=0.76; 95% CI, 0.63–0.91; I<sup>2</sup>=21%; p=0.004) subgroups, which was similar with a previous meta-analysis that included nine studies (35). In the Per AF subgroup, the rate of AF recurrence was numerically lower in the CF group than in the NCF group; however, this did not reach statistical significance (49.7% vs. 55.8%; RR=0.93; 95% CI, 0.77–1.12; I<sup>2</sup>=53%; p=0.43; Fig. 3a).

Moreover, seven studies provided data on the rate of acute PV reconnection, and no evidence of heterogeneity was found among the studies ( $I^2=0\%$ ). The acute PV reconnection significantly improved using CF-sensing catheters compared with that using NCF-sensing catheters (10.1% vs. 24.2%; RR=0.45; 95% CI, 0.32–0.63;  $I^2=0\%$ ; p=0.00001; Fig. 3b).

The CF used in the included studies ranged between 10 and 40 g, and the mean CF used was 18.3 g.

#### Safety of AF ablation using CF-sensing catheters

As shown in Figure 4, 11 studies assessed the incidence rate of major complications, and no evidence of heterogeneity was found among these studies ( $I^2=0\%$ ). The incidence rate of major complications was significantly lower in the CF group than in the NCF group (1.8% vs. 3.1%; OR=0.59; 95% CI, 0.37–0.95;  $I^2=0\%$ ; p=0.03). The incidence rate of minor complications was numerically lower in the CF group than in the NCF group; however, the results did not reach statistical significance (5.4% vs. 5.8%; OR=1.22; 95% CI, 0.78–1.92;  $I^2=0\%$ ; p=0.37).

Most included studies provided data on procedure parameters such as procedure duration, ablation time, fluoroscopy duration, and radiation dose in the AF and PAF subgroups. Figure 5 show that in the AF subgroup, the procedure duration [MD=–28.35; 95% CI, -39.54 to -17.16; l<sup>2</sup>=85%; p=0.00001], ablation time(MD=–3.8; 95% CI, -6.6 to -1.0; l<sup>2</sup>=76%; p=0.008), fluoroscopy duration (MD=–8.18; 95% CI, -14.11 to -2.24; l<sup>2</sup>=97%; p=0.007), and radiation dose (SMD=–0.75; 95% CI, -1.32 to -0.18; l<sup>2</sup>=90%; p=0.01) significantly reduced in the CF-guided group compared with in the NCF group. In the PAF subgroup, the procedure duration (MD=–49.64; 95% CI, -76.5 to -22.78; l<sup>2</sup>=83%; p=0.0003), ablation time (MD=–4.68; 95% CI, -13.83 to -3.52; l<sup>2</sup>=67%; p=0.001), fluoroscopy duration (MD=–13.9; 95% CI, -22.25 to -5.55; l<sup>2</sup>=93%; p=0.0001), and radiation dose (SMD=–0.56; 95% CI, -1.04 to -0.08; l<sup>2</sup>=73%; p=0.02) significantly reduced in the CF-guided group compared with in the NCF group.

## Discussion

This meta-analysis showed that in contrast to AF and PAF ablation performed using NCF-sensing catheters, the use of CFsensing catheters resulted in a significantly lower rate of acute PV reconnection and AF recurrence during the 12-month followup as well as reduced major complications and procedure parameters related to safety.

	Contact force-guided a	blation	Standard radiofrequency a	blation		Odds ratio		Odds ratio
Study or subgroup	Events	Total	Events	Total	Weight	M–H, Fixed, 95% Cl	Year	M–H, Fixed, 95% Cl
2.1.1 Major complication	ons							
Martinek 2012	1	25	1	25	1.2%	1.00 [0.06, 16.93]	2012	
larman 2014	2	200	10	400	7.9%	0.39 [0.09, 1.82]	2014	
Akca 2014	3	143	24	306	18.0%	0.25 [0.07, 0.85]	2014	<b>_</b>
Marijon 2014	2	30	1	30	1.1%	2.07 [0.18, 24.15]	2014	
Vakili 2014	1	32	1	35	1.1%	1.10 [0.07, 18.29]	2014	
Vutzler 2014	0	31	1	112	0.8%	11.18 [0.05, 29.68]	2014	
Jllah 2014	2	50	2	50	2.3%	1.00 [0.14, 7.39]	2014	
Vakamura 2015	1	60	0	60	0.6%	3.05 [0.12, 76.39]	2015	
Reddy 2015	1	152	2	143	2.5%	0.47 [0.04, 5.21]	2015	
G. Lee 2015	9	510	25	1005	19.9%	0.70 [0.33, 1.52]	2015	_ <b>_</b>
Sigmund 2015	2	99	3	99	3.5%	0.66 (0.11, 4.04]	2015	
Subtotal (95% CI)		1332		2265	58.9%	0.59 [0.37, 0.95]		$\bullet$
otal events	24		70					
leterogereity: Chi <sup>2</sup> =5.16	6, df=10 (P=0.88); l²=0%							
est for overall effect: Z								
.1.2 Minor complication	ons							
/lartinek 2012	1	25	3	25	3.5%	0.31 [0.03, 3.16]	2012	
laldar 2012	1	20	0	20	0.6%	3.15 [0.12, 82.16]	2012	
Varijon 2014	1	30	2	30	2.3%	0.48 [0.04, 5.63]	2014	
Jllah 2014	1	50	0	50	0.6%	3.6 [0.12, 76.95]	2014	
Vutzer 2014	1	31	3	112	1.5%	1.21 [0.12, 12.7]	2014	
Vakili 2014	1	32	1	35	1.1%	1.10 [0.07, 18.29]	2014	
Akca 2014	24	143	37	306	23.6%	1.47 [0.84, 2.56]	2014	+=-
Volf 2015	1	24	0	12	0.7%	1.60 [0.06, 42.13]	2015	
lakamura 2015	1	60	1	60	1.2%	1.00 [0.06, 16.37]	2015	
igmund 2015	1	99	2	99	2.4%	0.49 [0.04, 5.55]	2015	
Reddy 2015	3	152	3	143	3.6%	0.94 [0.19, 4.73]	2015	
Subtotal (95% CI)		666		892	41.1%	1.22 [0.78, 1.92]	2015	◆
otal events	36		52					
leterogereity: Chi <sup>2</sup> =3.64	4, df=10 (P=0.96); l <sup>2</sup> =0%							
est for overall effect: Z	Z=0.89 (P=0.37)							
otal (95% CI)		1998		3157	100.0%	085 [0.62, 1.17]		•
otal events	60		122					
	89, df=21 (P=0.91); l <sup>2</sup> =0%							0.002 0.1 1 10 5
est for overall effect: Z								Favours [experimental] Favours [control]
est for subgroup differ	rences: Chi <sup>2</sup> =4.80, df=1 (P=	=0.03); l <sup>2</sup> =79.2	%					ravours (experimental) ravours (control)

Figure 4. Forest plot showing odds ratio and 95% CI for the incidence rate of major complications and minor complications for studies comparing the CF and NCF groups

Achieving a lasting conduction block during the ablation procedure depends on a multitude of factors, including tissue depth, electrode-tissue interface temperature, and electrode tip-tissue contact pressure (29). Insufficient CF during initial lesion formation may result in edema and ineffective non-transmural lesions that allow subacute PV reconnection when the edema resolves (2, 12), whereas excessive contact can cause collateral tissue injury (31, 32, 36). Conventionally, the adequacy of contact between a catheter tip and tissue has been assessed using a combination of subjective factors and objective ablation parameters. Unfortunately, these parameters are poor predictors as they are unreliable and difficult to use (29, 37).

CF-sensing catheters offer a new paradigm in the invasive management of AF. Using these, continuous catheter-tissue CF can be measured, which ensures not only the optimal initial placement of the catheter but also the ability to detect catheter dislodging/sliding in real time (31). According to these features, the use of CF technology resulted in a significant reduction in the rate of acute PV reconnection and AF recurrence after AF ablation compared with the use of NCF.

However, it is a challenge to identify the optimal CF that should be applied during AF ablation to ensure adequate lesion formation, avoiding collateral tissue injury by the mean time.

The TOCCATA study (38) demonstrated that when PV isolation was performed with an average CF of <10 g, AF recurrence was

100%. When the average CF was >20 g, AF recurrence reduced to 20%. A recent published study (39) demonstrated that a CF threshold of >12 g predicts a complete lesion with high specificity. In the TOCCASTAR study, Reddy et al. (16) demonstrated that ablation with an optimal CF (≥90% of lesions created with a CF of  $\geq$ 10 g) resulted in a significantly higher success rate than that obtained for PV isolation with a non-optimal CF. The EFFICAS II study (40) prospectively applied CF guidelines for ensuring durable isolation of the PV of PAF patients, which demonstrated a target CF of 20 g; a range of 10-30 g resulted in a superior rate of durable PV isolation than the similar protocol without guidelines. The SMART-AF trial, a prospective, multicenter, non-randomized study (41), demonstrated that with an average CF of 17.9±9.4 g, 72.5% of patients were free from AF recurrence in a 12-month follow-up. The current meta-analysis provided important information regarding the use of an optimal average CF of 18.3 g (range, 10–40 g), with acceptable recurrence and complication rates.

Whether the use of CF-sensing catheters can decrease the rate of complications after AF ablation has always been a controversial issue. Akça et al. (32) demonstrated that CF procedures are associated with lesser major complications during AF ablation than NCF ones (2.1% vs. 7.8%, p=0.01). A previous metaanalysis (42) that included 11 studies demonstrated that the major complication rate was numerically lower in the CF group than in the NCF group; however, this did not reach statistical signifi-

а	Contact for	ce-guided a	blation	Standard ra	diofrequency a	ablation		Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	ŠD .	Total	Weight	IV, Random, 95% CI Year	IV, Random, 95% CI
2.2.1 AF									
Haldar 2012	209	65	20	207	59	20	4.3%	2.00 [-36.47, 40.47] 2012	
Martinek 2012	154	39	25	185	46	25	6.2%	-31.00 [-54.64, -7.36] 2012	
Sciarra 2014	140	53	21	181	53	21	5.0%	-41.00 [-73.06, -8.94] 2014	
Kimura 2014	59	16	19	96	39	19	6.9%	-37.00 [-55.95, -18.05] 2014	
Akca2014	191	56	143	194	72	306	7.8%	-3.00 [-15.22, 9.22] 2014	
Wutzler 2014	128.4	29	31	157.7	30.8	112	7.8%	-29.30 [-40.99, -17.61] 2014	<u> </u>
Wakili 2014	78.1	7.2	32	95.5	7.4	35	8.5%	-17.40 [-20.90, -13.90] 2014	-
Sigmund 2015	192	53	99	226	53	99	7.4%	-34.00 [-48.76, -19.24] 2015	_ <b>_</b>
Itoh 2015	160	30	50	245	61	50	6.9%	-85.00 [-103.84, -66.16] 2015	_ <b>_</b>
Wolf 2015	117.9	23.3	24	134.1	25.3	12	7.1%	-16.20 [-33.28, 0.88] 2015	
Makimoto 2015	133	42	35	152	33	35	7.0%	-19.00 [-36.70 -1.30] 2015	
Subtotal (95% CI)			499			734	74.9%	-28.35 [-39.54, -17.16]	◆
Heterogeneity: Tau <sup>2</sup> =2 Test for overall effect			0.00001); I²=	-85%					
2.2.2 Paroxysmal AF									
Martinek 2012	154	39	25	185	46	25	6.2%	-31.00 [-54.64, -7.36] 2012	
Sciarra 2014	140	53	21	181	53	21	5.0%	-41.00 [-73.06, -8.94] 2014	
Sigmund 2015	178.3	50.7	62	216.9	54	64	7.0%	-38.60 [-56.88, -20.32] 2015	
Itoh 2015	160	30	50	245	61	50	6.9%	-85.00 [-103.84, -66.16] 2015	
Subtotal (95% CI)	00.40.01.2.47.0		158	<b>N</b> 0/		160	25.1%	-49.64 [-76.50, -22.78]	
Heterogeneity: Tau <sup>2</sup> =6 Test for overall effect			.0006); 1²=83	5%					
Total (95% CI)			657			894	100.0%	-33.84 [-45.10, -22.59]	•
Heterogeneity: Tau <sup>2</sup> =3	386.44: Chi <sup>2</sup> =117.	12. df=14 (P		<sup>2</sup> =88%					
Test for overall effect									
Test for subgroup diff			0.15),   <sup>2</sup> =51	4%					Favours [experimental] Favours [control]
0 1		,							
b	Contact for	ce-quided a	hlation	Standard ra	diofrequency a	ablation		Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	SD		Weight	IV, Random, 95% Cl Year	IV, Random, 95% Cl
221 AE		55			55	. 5141	giit	,,,	,

	Contact forc	e-guided a:	iblation	Standard ra	diofrequency	ablation		Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI Yea	r IV, Random, 95% Cl
2.3.1 AF									
Martinek 2012	39	11	25	50.5	15.9	25	5.3%	-11.50 [-19.08, -3.92] 2012	
Haldar 2012	60.7	20.6	20	51.9	23.1	20	2.7%	8.80 [-4.76, 22.36] 2012	
Wakili 2014	30.8	3.9	32	31.7	2.4	35	9.3%	-0.90 [-2.47, 0.67] 2014	
Sciarra 2014	30	14	21	41.3	13.2	21	4.9%	-11.30 [-19.53, -3.07] 2014	
Wutzler 2014	38.6	12.7	31	45.2	16.5	112	6.8%	-6.60 [-12.02, -1.18] 2014	
Andrade 2014	58.8	22.1	25	56.4	24	50	3.6%	2.40 [-8.52, 13.32] 2014	
Jarman 2014	55	23	200	54	24	400	7.9%	1.00 [-2.96, 4.96] 2014	
Marijon 2014	45.2	18	30	65.4	22	30	3.9	-20.20 [-30.37, 1-10.03] 2014	
Wolf 2015	31.5	7.1	24	31.8	7	12	7.2%	-0.30 [-5.17, 4.57] 2015	
Makimoto 2015	13.1	3.6	35	13.2	4.3	35	9.1%	-0.10 [-1.96, 1.76] 2015	+
Sigmund 2015	43.6	16.4	99	51.8	19.6	99	7.1%	-8.20 [-13.23, -3.17] 2015	
Subtotal (95% CI)	00 01 3 40 54	16 40 /D 0	542	n/		839	67.8%	-3.80 [-6.60, -1.00]	•
Heterogeneity: Tau <sup>2</sup> =12			.00001); 1²=/b	%					
Test for overall effect: 2	2=2.66 (P=0.008	i)							
2.3.2 Paroxysmal AF									
Martinek 2012	39	11	25	50.5	15.9	25	5.3%	-11.50 [-19.08, -3.92] 2012	
Marijon 2014	45.2	18	30	65.4	22	30	3.9%	-20.20 [-30.37, -10.03] 2014	
Andrade 2014	58.8	22.1	25	56.4	24	50	3.6%	2.40 [-8.52, 13.32] 2014	
Jarman 2014	45	16	92	48	22	184	7.4%	-3.00 [-7.56, 1.56] 2014	
Sciarra 2014	30	14	21	41.3	13.2	21	4.9%	-11.30 [-19.53, -3.07] 2014	
Sigmund 2015	38.5	12.9	62	48.1	17	64	6.9%	-9.60 [-14.86, -4.34] 2015	
Subtotal (95% CI)	00.0	12.0	255			374	32.2%	-8.68 [-13.83, -3.52]	
Heterogeneity: Tau <sup>2</sup> =26	.35: Chi <sup>2</sup> =156.37	7. df=5 (P=0	.009): l <sup>2</sup> =67%						
Test for overall effect: 2									
		-							
Total (95% CI)			797			1213	100.0%	-5.47 [-8.10, -2.84]	◆
Heterogeneity: Tau <sup>2</sup> =18	.82; Chi <sup>2</sup> =75.25,	df=16 (P<0	.00001); l <sup>2</sup> =79	%					
Test for overall effect: 2									Favours [experimental] Favours [control]
Test for subgroup differ	rences: Chi <sup>2</sup> =2.	65, df=1 (P=	=0.10), l <sup>2</sup> =62.3	%					
L									

Figure 5. (a–c) Forest plot showing the unadjusted difference in the mean procedure duration, ablation time, and fluoroscopy duration for studies comparing the CF and NCF groups. (d) Forest plot showing the standard difference in the mean radiation dose for studies comparing the CF and NCF groups

cance (1.3% vs. 1.9%; OR, 0.71; 95% CI, 0.29–1.73; p=0.45). With more studies included, the current meta-analysis demonstrated that the incidence of major complications was significantly lower in the CF group than in the NCF group (1.8% vs. 3.1%; OR=0.59; 95% CI, 0.37–0.95; p<0.05).

In the current analysis, the procedure duration, ablation time, fluoroscopy duration, and radiation dose significantly reduced in the CF group compared with in the NCF group in the AF and PAF subgroups. CF-sensing catheters may reduce reliance on fluoroscopy during navigation and the time to achieve intact linear lesions, which promote safety not only for patients but also for operators.

#### **Study limitations**

The current analysis had the following limitations: some studies were of limited quality, given their retrospective and single-center designs. Differences in operators' experience and ablation protocols may have affected the outcomes of the included studies.

## Conclusion

AF ablation using CF-sensing catheters has better outcomes than those NCF-sensing catheters during the 12-month followup period. Furthermore, the incidence of major complications

C		e-guided abl		Standard radi				Mean difference	Mean difference
tudy or subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	IV, Random, 95% Cl
4.1 AF									
kca 2014	57.5	20.1	143	45.7	24.2	306	6.4%	11.80 [7.53, 16.07]	
oh 2015	17	8	50	54	27	50	5.7%	-37.00 [-44.81, -29.19]	
arman 2014	26.6	15.1	92	34.7	18.7	184	6.4%	-8.10 [-12.20, -4.00]	-
imura 2004	9	20	19	22	63	19	1.8%	-13.00 [-42.72, 16.72]]	
Aakimoto 2015	13.5	6.6	35	15.7	6.5	35	6.6%	-2.20 [-5.27, 0.87]	-
larijon 2014	20.1	4	30	26.7	5	30	6.6%	-6.60 [-8.89, -4.31]	<b>-</b>
Aartinek 2012	23.6	13.1	25	28.6	17.4	25	5.5%	-5.00 [-13.54, 3.54]	_ <b>_</b> +
ciarra 2014	20.0	10	21	34	18	21	5.4%	-14.00 [-22.81, -5.19]	
igmund 2015	19.9	9.3	99	28.5	11	99	6.6%	-8.60 [-11.44, -5.76]	
Vakili 2014	33	2.7	33	51.4	3.3	35	6.7%	-18.40 [-19.84, -16.96]	-
							6.4%		<u> </u>
Volf 2015	11.8	5.6	24	11	5.8	12		0.80 [-3.17, 4.77]	
Vutzler 2014	39.7	11.3	31	43.8	14.5	112	6.3%	-4.10 [-8.90, 0.70]	<b>A</b>
ubtotal (95% CI)			601			928	<b>70.3</b> %	-8.18 [-14.11, -2.24]	◆
eterogeneity: Tau <sup>2</sup> =96 est for overall effect:			)0001); l²=97	%					
4.2 Paroxysmal AF									
oh 2015	17	8	50	54	27	50	5.7%	-37.00 [-44.81, -29.19]	——
Aarijon 2014	20.1	4	30	26.7	5	30	6.6%	-6.60 [-8.89, -4.31]	-
Aartinek 2012	23.6	13.1	25	28.6	17.4	25	5.5%	-5.00 [-13.54, 3.54]	+
ciarra 2014	20	10	21	34	18	21	5.4%	-14.00 [-22.81, -5.19]	
iamund 2015	18.6	8.8	62	27.3	12.1	64	6.5%	-8.70 [-12.39, -5.01]	
ubtotal (95% CI)	10.0	0.0	188	27.5	12.1	190	<b>29.7%</b>	-13.90 [-22.25, -5.55]	◆
leteroneneuty. Taur=75	9 58 1.01 = 55 / 5								
est for overall effect:	Z=3.26 (P=0.001	)	789	0/		1118	100.0%	-9.84 [-14.45, -5.23]	•
est for overall effect: otal leterogeneity: Tau <sup>2</sup> =81 est for overall effect:	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.35 Z=4.18 (P<0.000	) 5, df=16 (P<0.0 11)	<b>789</b> 00001); l²=96			1118	100.0%	-9.84 [-14.45, -5.23]	← 100 -50 50 Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau2=81 est for overall effect: est for subgroup diffe tudy or subgroup	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.35 Z=4.18 (P<0.000 prences: Chi <sup>2</sup> =1.	) 5, df=16 (P<0.0 11)	<b>789</b> 00001); l <sup>2</sup> =96 27), l <sup>2</sup> =16.6%		ofrequency ab SD	lation	100.0% Weight	–9.84 [–14.45, –5.23] Std. Mean difference IV, Random, 95% Cl	
est for overall effect: otal eterogeneity: Tau2=81 est for overall effect: est for subgroup diffe tudy or subgroup 5.1 AF	Z=3.26 (P=0.001 1.87; Chi²=405.38 Z=4.18 (P<0.000 rences: Chi²=1. Contact forc <u>Mean</u>	) 5, df=16 (P<0.( 11) 20, df=1 (P=0. 20, df=1 ( <b>P=0</b> . <b>20</b> <b>SD</b>	789 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% ation Total	Standard radio Mean	SD	lation Total	Weight	Std. Mean difference IV, Random, 95% Cl	Favours [experimental] Favours [control]
est for overall effect: . total leterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup 5.1 AF casella 2014	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rrences: Chi <sup>2</sup> =1.: Contact forc <u>Mean</u> 2.069	) 5, df=16 (P<0.( 11) 20, df=1 (P=0. 20, df=1 ( <b>P=0.</b> <b>SD</b> 649	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <u>Total</u> 20	Standard radio Mean 2.030	695	lation Total 35	Weight 8.9%	Std. Mean difference IV, Random, 95% CI 0.06 [–0.49, 0.61]	Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup .5.1 AF asella 2014 Aakimoto 2015	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1. Contact forc <u>Mean</u> 2.069 2.047	) 5, df=16 (P<0.( 11) 20, df=1 (P=0. <b>:e-guided abl</b> <b>SD</b> 649 973	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% ation Total 20 35	Standard radio Mean 2.030 2.281	695 1.229	lation Total 35 35	<b>Weight</b> 8.9% 9.3%	Std. Mean difference IV, Random, 95% Cl 0.06 [-0.49, 0.61] -0.21 [-0.68, 0.26]	Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau²=81 est for overall effect: est for subgroup diffe dudy or subgroup 5.1 AF easella 2014 Aakimoto 2015 Aarijon 2014	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1.: Contact forc <u>Mean</u> 2.069 2.047 41.6	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>:e-guided abl</b> <b>SD</b> 649 973 10	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30	Standard radio Mean 2.030 2.281 56.7	695 1.229 14	lation Total 35 35 30	Weight 8.9% 9.3% 8.8%	Std. Mean difference IV, Random, 95% CI 0.06 [-0.49, 0.61] -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67]	Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup .5.1 AF asella 2014 Aakinoto 2015 Martinek 2012	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rrences: Chi <sup>2</sup> =1.3 Contact forc <u>Mean</u> 2.069 2.047 41.6 58.510	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>:e-guided abl</b> <b>SD</b> 649 973 10 14.655	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30 25	5 <b>Standard radid</b> Mean 2.030 2.281 56.7 70.926	695 1.229 14 19.470	lation Total 35 35 30 25	Weight 8.9% 9.3% 8.8% 8.7%	Std. Mean difference IV, Random, 95% CI 0.06 [-0.49, 0.61] -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67] -0.71 [-1.28, -0.14]	Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup 5.1 AF asella 2014 fariinot 2015 fartinek 2012 igmund 2015	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1.3 Contact forc <u>Mean</u> 2.069 2.047 41.6 58.510 56.7	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>ce-guided abl</b> <b>SD</b> 649 973 10 14.655 38.9	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30 25 99	5 <b>Standard radi</b> Mean 2.030 2.281 56.7 70.926 74.1	695 1.229 14 19.470 58	lation Total 35 35 30 25 99	Weight 8.9% 9.3% 8.8% 8.7% 10.2%	Std. Mean difference IV, Random, 95% CI -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67] -0.71 [-1.28, -0.7] -0.35 [-0.63, -0.07]	Favours [experimental] Favours [control]
est for overall effect: otal eterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup 5.1 AF asella 2014 1akimoto 2015 1arijon 2014 1artinek 2012 igmund 2015 Vakili 2015	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1 Contact forc <u>Mean</u> 2.069 2.047 41.6 58.510 56.7 34.1	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>:e-guided abl</b> <b>SD</b> 649 973 10 14.655 38.9 3.1	<b>789</b> 00001}; I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30 25 99 32	<b>Standard radii</b> Mean 2.030 2.281 56.7 70.926 74.1 44.2	695 1.229 14 19.470 58 3.7	lation Total 35 35 30 25 99 35	Weight 8.9% 9.3% 8.8% 8.7% 10.2% 8.0%	Std. Mean difference IV, Random, 95% Cl 0.06 [-0.49, 0.61] -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67] -0.71 [-1.28, -0.14] -0.35 [-0.63, -0.07] -2.92 [-3.61, -2.21]	Favours [experimental] Favours [control]
est for overall effect: otal leterogeneity: Tau <sup>2</sup> =81 est for overall effect: est for subgroup diffe tudy or subgroup 5.1 AF asella 2014 Martinoto 2015 Marijon 2014 dartinek 2012 igmund 2015 Vakili 2014	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1.3 Contact forc <u>Mean</u> 2.069 2.047 41.6 58.510 56.7	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>ce-guided abl</b> <b>SD</b> 649 973 10 14.655 38.9	<b>789</b> 00001); I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30 25 99 32 31	5 <b>Standard radi</b> Mean 2.030 2.281 56.7 70.926 74.1	695 1.229 14 19.470 58	lation Total 35 35 30 25 99 35 112	Weight 8.9% 9.3% 8.8% 8.7% 10.2% 8.0% 9.7%	Std. Mean difference IV, Random, 95% CI -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67] -0.71 [-1.28, -0.7] -0.35 [-0.63, -0.07]	Favours [experimental] Favours [control]
ast for overall effect: otal eterogeneity: Tau <sup>2</sup> =81 est for overall effect: ast for subgroup diffe tudy or subgroup 5.1 AF asella 2014 larijon 2015 larijon 2014 lartinek 2012 igmund 2015 Jakili 2014 Jutzler 2014	Z=3.26 (P=0.001 1.87; Chi <sup>2</sup> =405.38 Z=4.18 (P<0.000 rences: Chi <sup>2</sup> =1 Contact forc <u>Mean</u> 2.069 2.047 41.6 58.510 56.7 34.1	) 5, df=16 (P<0.( 11) 20, df=1 (P=0.) <b>:e-guided abl</b> <b>SD</b> 649 973 10 14.655 38.9 3.1	<b>789</b> 00001}; I <sup>2</sup> =96 27), I <sup>2</sup> =16.6% <b>ation</b> <b>Total</b> 20 35 30 25 99 32	<b>Standard radii</b> Mean 2.030 2.281 56.7 70.926 74.1 44.2	695 1.229 14 19.470 58 3.7	lation Total 35 35 30 25 99 35	Weight 8.9% 9.3% 8.8% 8.7% 10.2% 8.0%	Std. Mean difference IV, Random, 95% Cl 0.06 [-0.49, 0.61] -0.21 [-0.68, 0.26] -1.23 [-1.78, -0.67] -0.71 [-1.28, -0.14] -0.35 [-0.63, -0.07] -2.92 [-3.61, -2.21]	Favours [experimental] Favours [control]
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Heterogeneity: Tau<sup>2</sup>=0.36; Chi<sup>2</sup>=73.33, df=10 (P<0.00001); l<sup>2</sup>=86% Test for overall effect: Z=3.43 (P=0.0006) Test for subgroup difference: Chi<sup>2</sup>=0.24, df=1 (P=0.62), l<sup>2</sup>=0%

Continued Figure 5. (a–c) Forest plot showing the unadjusted difference in the mean procedure duration, ablation time, and fluoroscopy duration for studies comparing the CF and NCF groups. (d) Forest plot showing the standard difference in the mean radiation dose for studies comparing the CF and NCF groups.

using CF-sensing catheters was even lower than that using NCF-sensing catheters. The meta-analysis also demonstrated that using an optimal average CF of 18.3 g was associated with higher success and lower complication rates. Randomized controlled studies are required to assess whether catheter ablation using an optimized CF improves the long-term clinical outcome and to determine the exact optimal CF to be used in different patient subgroups.

## Conflict of interest: None declared.

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Favours [experimental] Favours [control]

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