

# **Role katetrizační ablace fibrilace síní v léčbě srdečního selhání**

**Jan Škoda**

**Nemocnice Na Homolce**



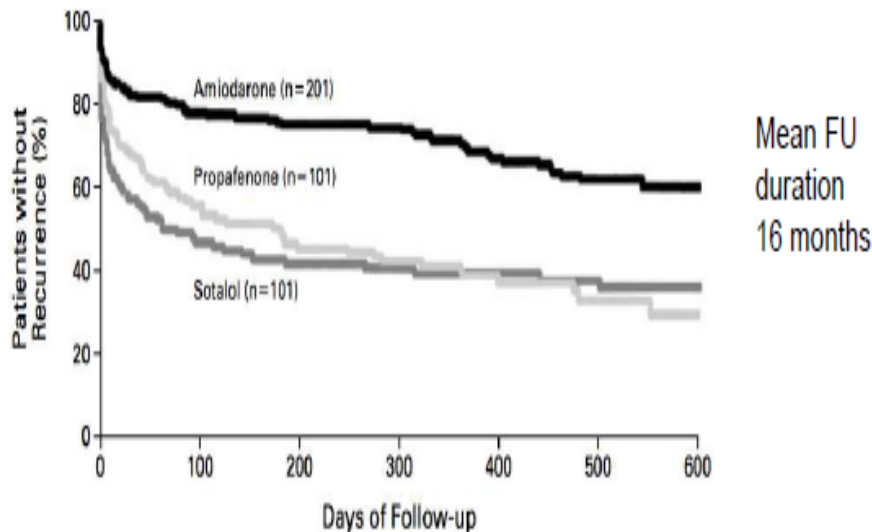
# Co očekáváme od katetrizační ablace FiS u srdečního selhání

- Zlepšení klinických obtíží
- Zlepšení tolerance zátěže
- Zlepšení kvality života
- Snížení rizika CMP
- Zlepšení ejekční frakce LK
- Mortalitní benefit

# Limitace účinku antiarytmik

## AMIODARONE TO PREVENT RECURRENCE OF ATRIAL FIBRILLATION

DENIS ROY, M.D., MARIO TALAJIC, M.D., PAUL DORIAN, M.D., STUART CONNOLLY, M.D.,  
 MARK J. EISENBERG, M.D., M.P.H., MARTIN GREEN, M.D., TERESA KUS, M.D., JEAN LAMBERT, PH.D.,  
 MARC DUBUC, M.D., PIERRE GAGNÉ, M.D., STANLEY NATTEL, M.D., AND BERNARD THIBAUT, M.D.,  
 FOR THE CANADIAN TRIAL OF ATRIAL FIBRILLATION INVESTIGATORS\*



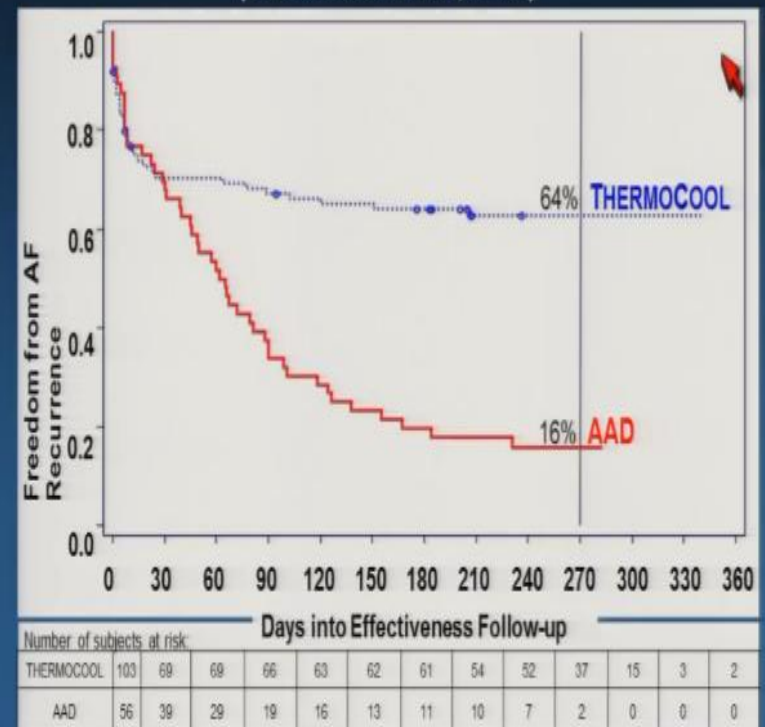
AF Recurrence 35% with Amiodarone and 63% with sotalol and propafenone (P<0.001)

Roy et al. NEJM 2000

## NAVISTAR® THERMOCOOL® Catheter for the Radiofrequency Ablation of Symptomatic Paroxysmal Atrial Fibrillation

PMA # P030031/S11

KM Curve of Time to Chronic Failures By Randomization Group  
 (Effectiveness Cohort, n=159)



	Days into Effectiveness Follow-up												
	0	30	60	90	120	150	180	210	240	270	300	330	360
Number of subjects at risk													
THERMOCOOL	103	69	69	66	63	62	61	54	52	37	15	3	2
AAD	56	39	29	19	16	13	11	10	7	2	0	0	0

# Limitace účinku antiarytmik

ORIGINAL ARTICLE

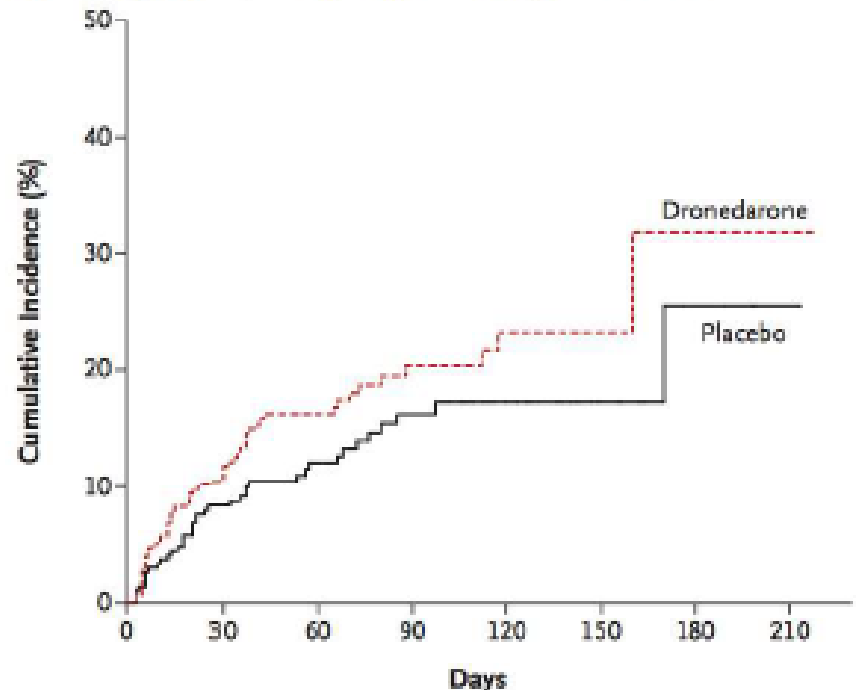
## Increased Mortality after Dronedarone Therapy for Severe Heart Failure

Lars Køber, M.D., Christian Torp-Pedersen, M.D., John J.V. McMurray, M.D., Ole Gøtzsche, M.D., Samuel Lévy, M.D., Harry Crijns, M.D., Jan Amle, M.D., and Jan Carlsson, M.D., for the Dronedarone Study Group<sup>†</sup>

- **Multicenter study with a double-blind design**
- **Planned to randomly assign 1000 patients who were hospitalized with symptomatic heart failure and severe left ventricular systolic dysfunction to receive 400 mg of dronedarone twice a day or placebo**
- **The primary end point was the composite of death from any cause or hospitalization for heart failure**
- **Study was terminated earlier after higher rate of death in the Dronedarone group**

Kober et al NEJM 2008

A All-Cause Mortality or Hospitalization for Worsening Heart Failure



No. at Risk

Placebo	317	234	159	87	41	16	6	1
Dronedarone	310	232	151	87	49	19	4	1

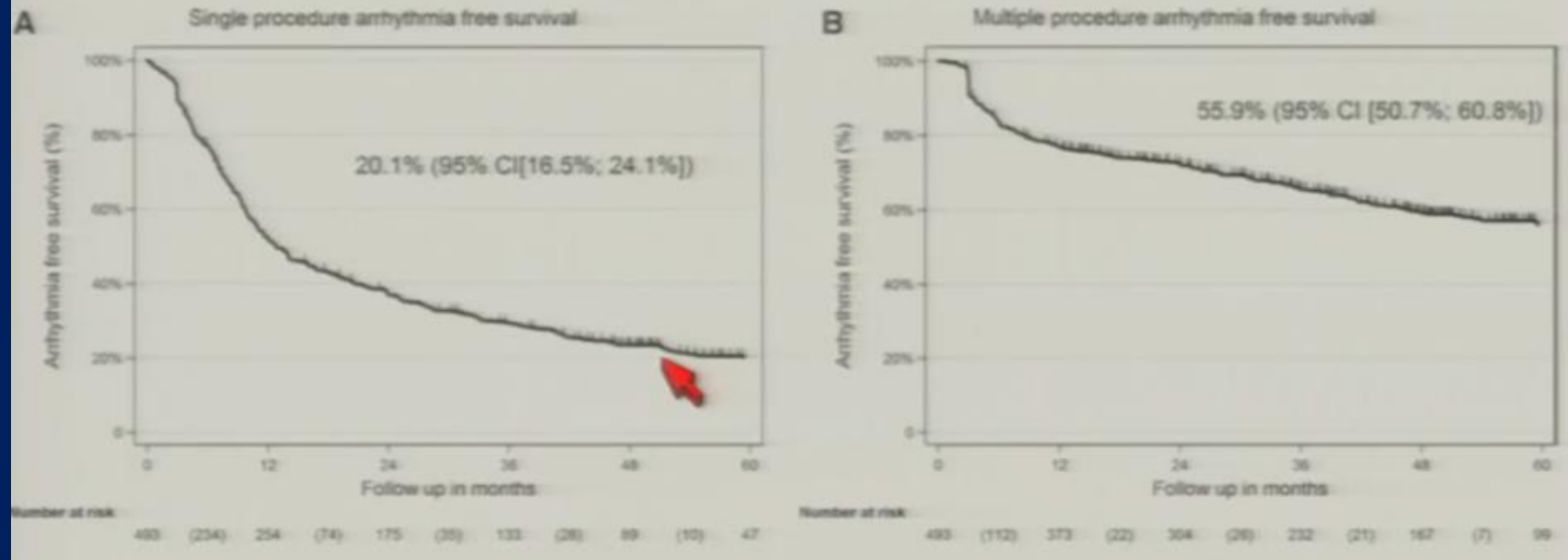


# Dlouhodobá efektivita katetrizační ablace FiS

## Persistent Atrial Fibrillation Using the Stepwise Approach and Prognostic Factors for Success

Doreen Schreiber, MD\*; Thomas Rostock, MD\*; Max Fröhlich; Arian Sultan, MD; Helge Servatius, MD; Boris A. Hoffmann, MD; Jakob Lüker, MD; Imke Berner, MD; Benjamin Schäffer, MD; Karl Wegscheider, PhD; Susanne Lezius, MSc; Stephan Willems, MD; Daniel Steven, MD

Circ A EP 2015;8:308-317.



# Izolace plicních žil vs ablace AVN u HF

THE NEW ENGLAND JOURNAL OF MEDICINE

ORIGINAL ARTICLE

## Pulmonary-Vein Isolation for Atrial Fibrillation in Patients with Heart Failure

Mohammed N. Khan, M.D., Feroz Jalil, M.D., Jennifer Cummings, M.D., Jang Di Saez, M.D., Prashanthan Sanderan, M.D., David G. Marmor, M.D., Josef Kostner, M.D., Steven Haas, M.D., Sotiris Themistoclakis, M.D., Raffaele Ferrari, M.D., Domenico Pistono, M.D., Maurizio Mazzoni, M.D., Ottaviano Vanni, M.D., Robert Schwartel, M.D., Waleed Salata, M.D., Paul Wang, M.D., Anna Pi-Ahmed, M.D., Satish Babunny, M.D., Pietro Santarelli, M.D., Rishi C. Starling, M.D., Antonio DeFina, M.D., Giovanni Pappalardo, M.D., Johannes Brachmann, M.D., Volker Schöngels, M.D., Aldo Bonato, M.D., Michela Cazzola, M.D., Rossana Boriani, M.D., Mohab Hammad, M.D., and Emelia Stehouwer, M.D., for the PAM-HF Investigators\*

ABSTRACT

**BACKGROUND** Pulmonary-vein isolation is increasingly being used to treat atrial fibrillation in patients with heart failure.

**OBJECTIVE** In this prospective, multicenter clinical trial, we randomly assigned patients with symptomatic, drug-resistant atrial fibrillation, an ejection fraction of 40% or less, and New York Heart Association class II or III heart failure to undergo either pulmonary-vein isolation or atrioventricular-node ablation with biventricular pacing. All patients completed the Minnesota Living with Heart Failure questionnaire (score range from 0 to 100, with a higher score indicating a worse quality of life) and underwent echocardiography and a 6-minute walk test the composite primary end point. Over a 6-month period, patients were monitored for both asymptomatic and symptomatic episodes of atrial fibrillation.

**RESULTS** In all, 41 patients underwent pulmonary-vein isolation, and 40 underwent atrioventricular-node ablation with biventricular pacing; none were lost to follow-up at 6 months. The composite primary end point favored the group that underwent pulmonary-vein isolation, with an improved questionnaire score at 6 months (66 vs. 82 in the group that underwent atrioventricular-node ablation with biventricular pacing;  $P=0.003$ ), a larger 6-minute-walk distance (340 m vs. 297 m,  $P=0.003$ ), and a higher ejection fraction (30% vs. 28%,  $P<0.001$ ) in the group that underwent pulmonary-vein isolation. 80% of patients receiving antiarrhythmic drugs and 73% of those not receiving such drugs were free of atrial fibrillation at 6 months. In the group that underwent pulmonary-vein isolation, pulmonary-vein stenosis developed in two patients, pericardial effusion in one, and pulmonary edema in another; in the group that underwent atrioventricular-node ablation with biventricular pacing, lead dislodgment was found in one patient and paroxysmal atrial fibrillation in another.

**CONCLUSION** Pulmonary-vein isolation was superior to atrioventricular-node ablation with biventricular pacing in patients with heart failure who had drug-resistant atrial fibrillation. (ClinicalTrials.gov number, NCT00999936.)

**INTRODUCTION** Pulmonary-vein isolation was superior to atrioventricular-node ablation with biventricular pacing in patients with heart failure who had drug-resistant atrial fibrillation. (ClinicalTrials.gov number, NCT00999936.)

From Cardiology Department, MRC Great Ormond Street Hospital, London, U.K. (Dr. Khan); Department of Cardiology, University of Liverpool, Liverpool, U.K. (Dr. Jalil); Cleveland Clinic, Cleveland, Ohio (Dr. Cummings); University of Liverpool, Liverpool, U.K. (Dr. Saez); University of Liverpool, Liverpool, U.K. (Dr. Sanderan); University of Liverpool, Liverpool, U.K. (Dr. Marmor); University of Liverpool, Liverpool, U.K. (Dr. Kostner); University of Liverpool, Liverpool, U.K. (Dr. Haas); University of Liverpool, Liverpool, U.K. (Dr. Themistoclakis); University of Liverpool, Liverpool, U.K. (Dr. Ferrari); University of Liverpool, Liverpool, U.K. (Dr. Pistono); University of Liverpool, Liverpool, U.K. (Dr. Vanni); University of Liverpool, Liverpool, U.K. (Dr. Schwartel); University of Liverpool, Liverpool, U.K. (Dr. Salata); University of Liverpool, Liverpool, U.K. (Dr. Wang); University of Liverpool, Liverpool, U.K. (Dr. Pi-Ahmed); University of Liverpool, Liverpool, U.K. (Dr. Babunny); University of Liverpool, Liverpool, U.K. (Dr. Santarelli); University of Liverpool, Liverpool, U.K. (Dr. Starling); University of Liverpool, Liverpool, U.K. (Dr. DeFina); University of Liverpool, Liverpool, U.K. (Dr. Pappalardo); University of Liverpool, Liverpool, U.K. (Dr. Brachmann); University of Liverpool, Liverpool, U.K. (Dr. Schöngels); University of Liverpool, Liverpool, U.K. (Dr. Bonato); University of Liverpool, Liverpool, U.K. (Dr. Cazzola); University of Liverpool, Liverpool, U.K. (Dr. Boriani); University of Liverpool, Liverpool, U.K. (Dr. Hammad); University of Liverpool, Liverpool, U.K. (Dr. Stehouwer).

\*The PAM-HF Investigators are: Mohammed N. Khan, M.D., Feroz Jalil, M.D., Jennifer Cummings, M.D., Jang Di Saez, M.D., Prashanthan Sanderan, M.D., David G. Marmor, M.D., Josef Kostner, M.D., Steven Haas, M.D., Sotiris Themistoclakis, M.D., Raffaele Ferrari, M.D., Domenico Pistono, M.D., Maurizio Mazzoni, M.D., Ottaviano Vanni, M.D., Robert Schwartel, M.D., Waleed Salata, M.D., Paul Wang, M.D., Anna Pi-Ahmed, M.D., Satish Babunny, M.D., Pietro Santarelli, M.D., Rishi C. Starling, M.D., Antonio DeFina, M.D., Giovanni Pappalardo, M.D., Johannes Brachmann, M.D., Volker Schöngels, M.D., Aldo Bonato, M.D., Michela Cazzola, M.D., Rossana Boriani, M.D., Mohab Hammad, M.D., and Emelia Stehouwer, M.D.

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N. Engl. J. Med. 2018;378:1718-28.

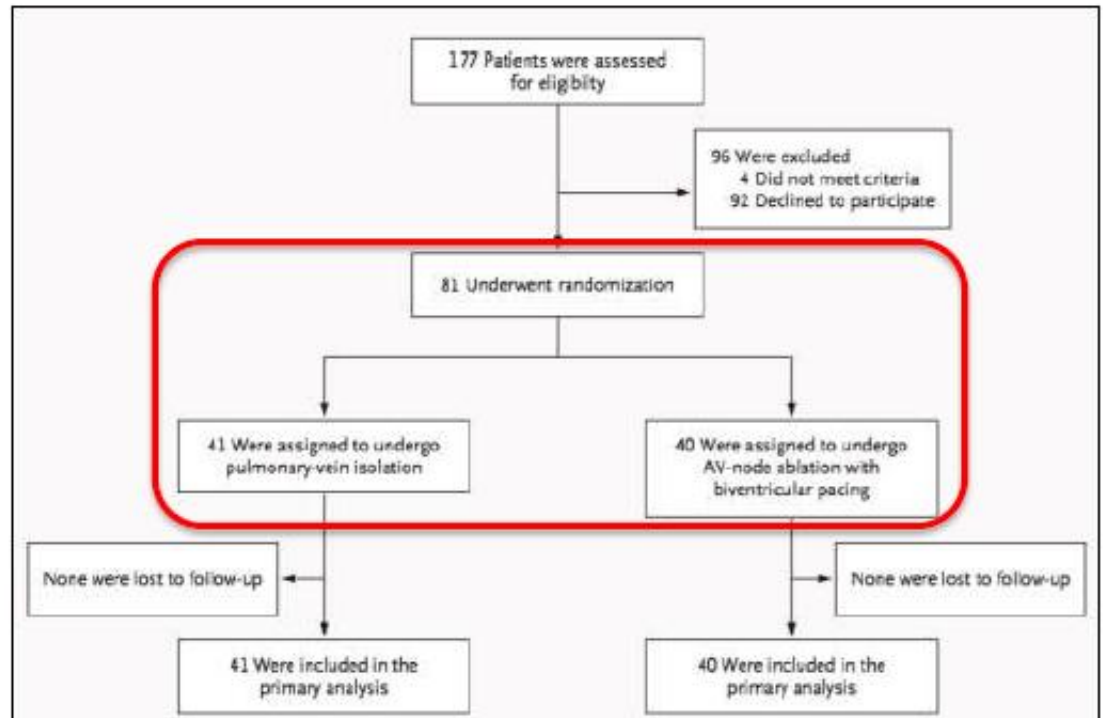
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DOI: 10.1056/NEJMoa1711718

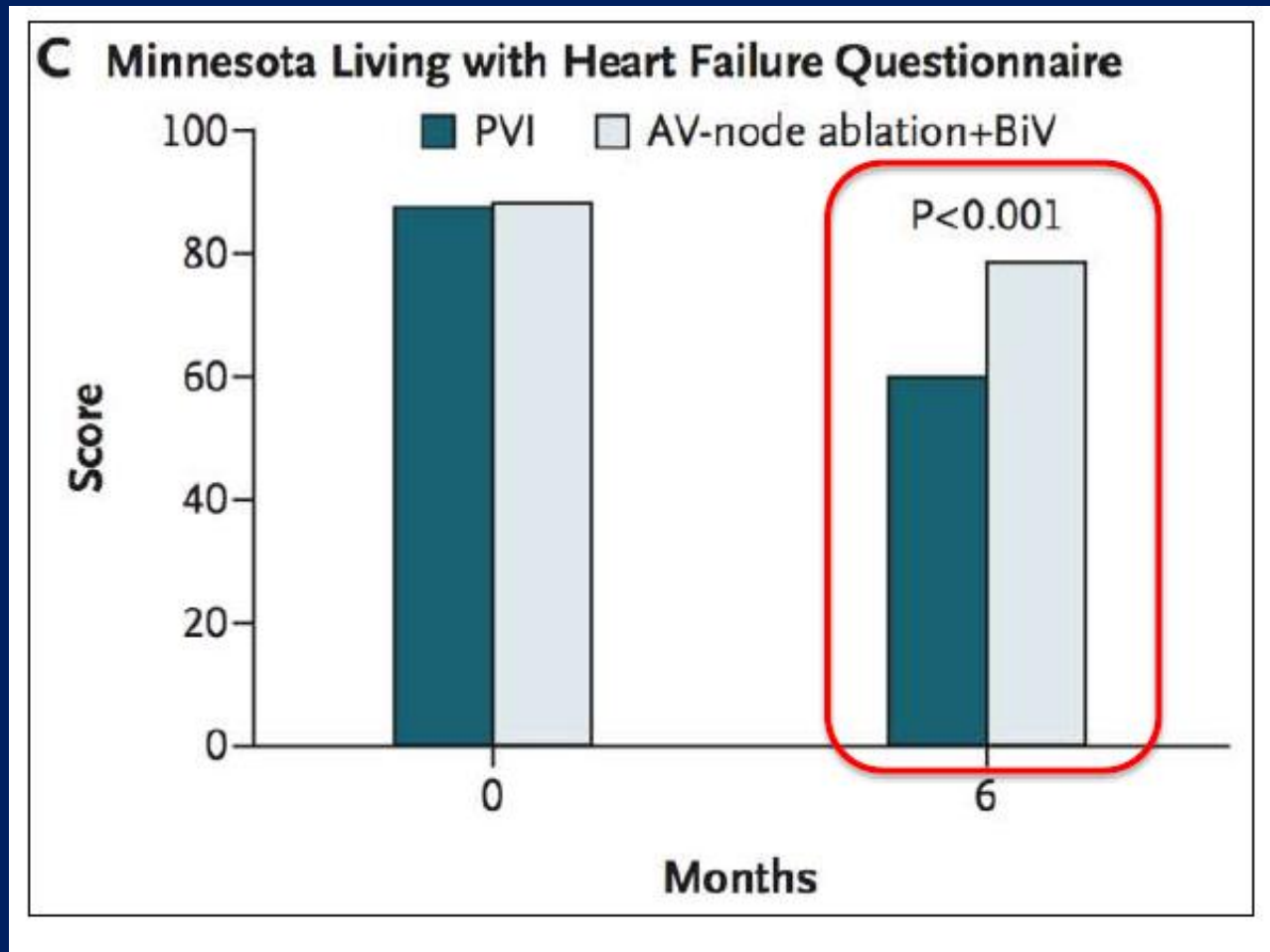
www.nejm.org

0007-1226/18/378-1718/\$18.00

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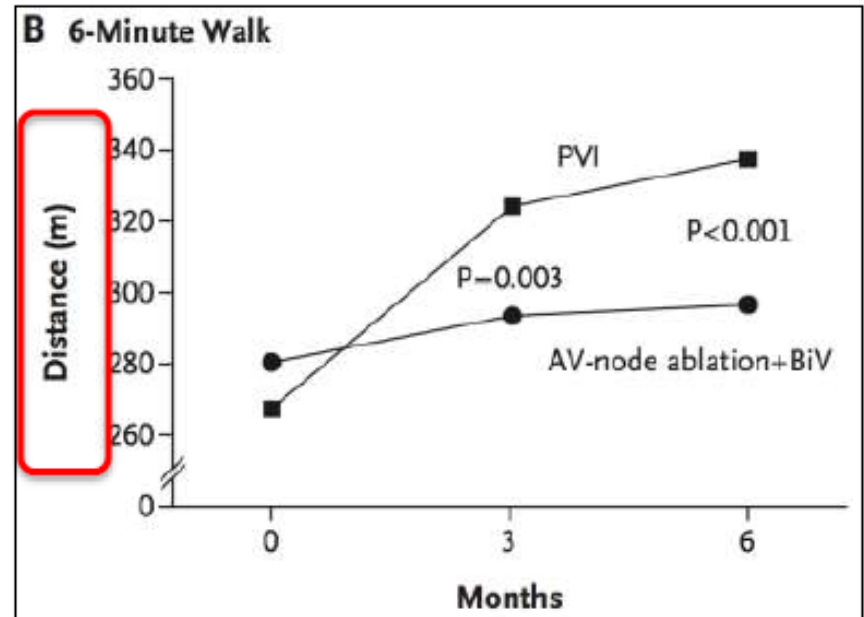
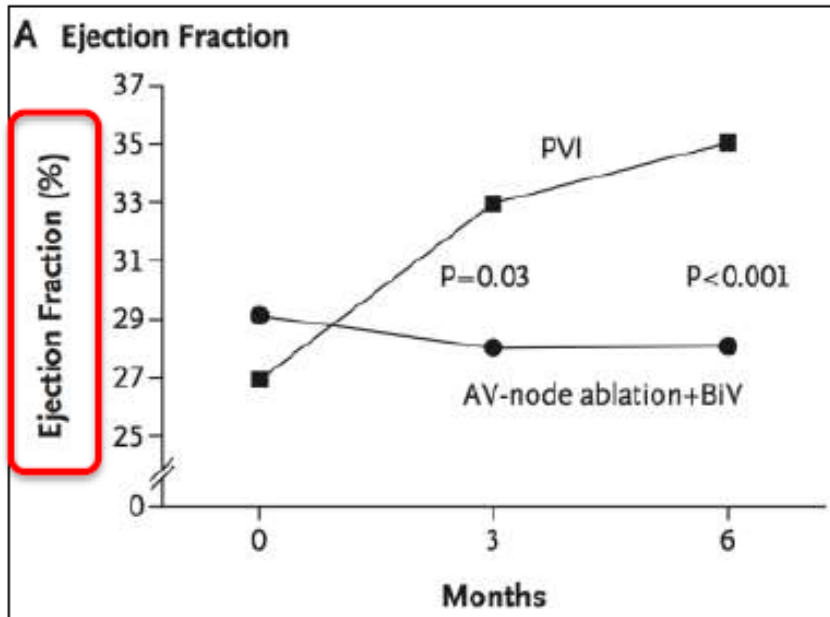


# Izolace plicních žil vs ablace AVN u HF





# Izolace plicních žil vs ablace AVN u HF



# Randomizovaná studie: ablace perzist FiS x rate control (ARC-HF)

## A Randomized Trial to Assess Catheter Ablation Versus Rate Control in the Management of Persistent Atrial Fibrillation in Heart Failure

David G. Jones, MD,\*† Shouvik K. Halder, MBBS,\*† Wajid Hussain, MB, CstB,\*† Rakesh Sharma, PhD,\*† Durol P. Francis, MD,† Shelley L. Rahman-Haley, MD,\* Theresa A. McDonagh, MD,\*† S. Richard Underwood, MD,\*† Vlas Markides, MD,\*† Tom Wong, MD\*†  
*London, United Kingdom*

EF pod 35%

FUP 12 mo

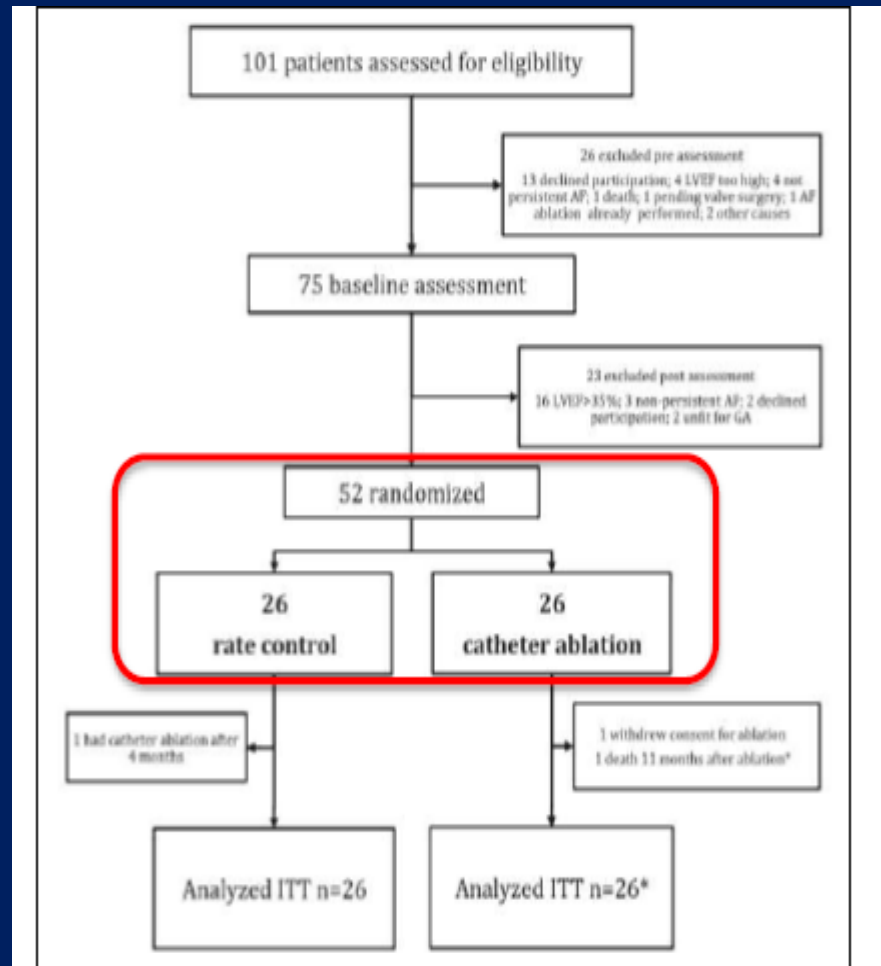
Primární cíl:  
změna Vo2 max

Sekund cíle:

BNP

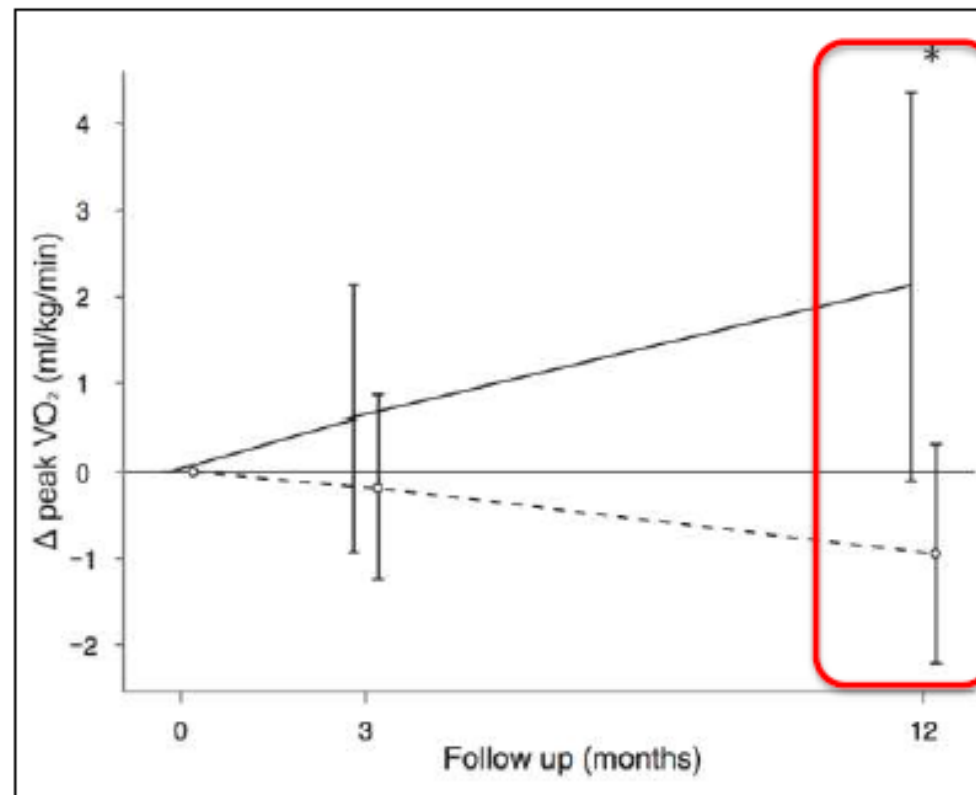
6min test chůze

EFLK



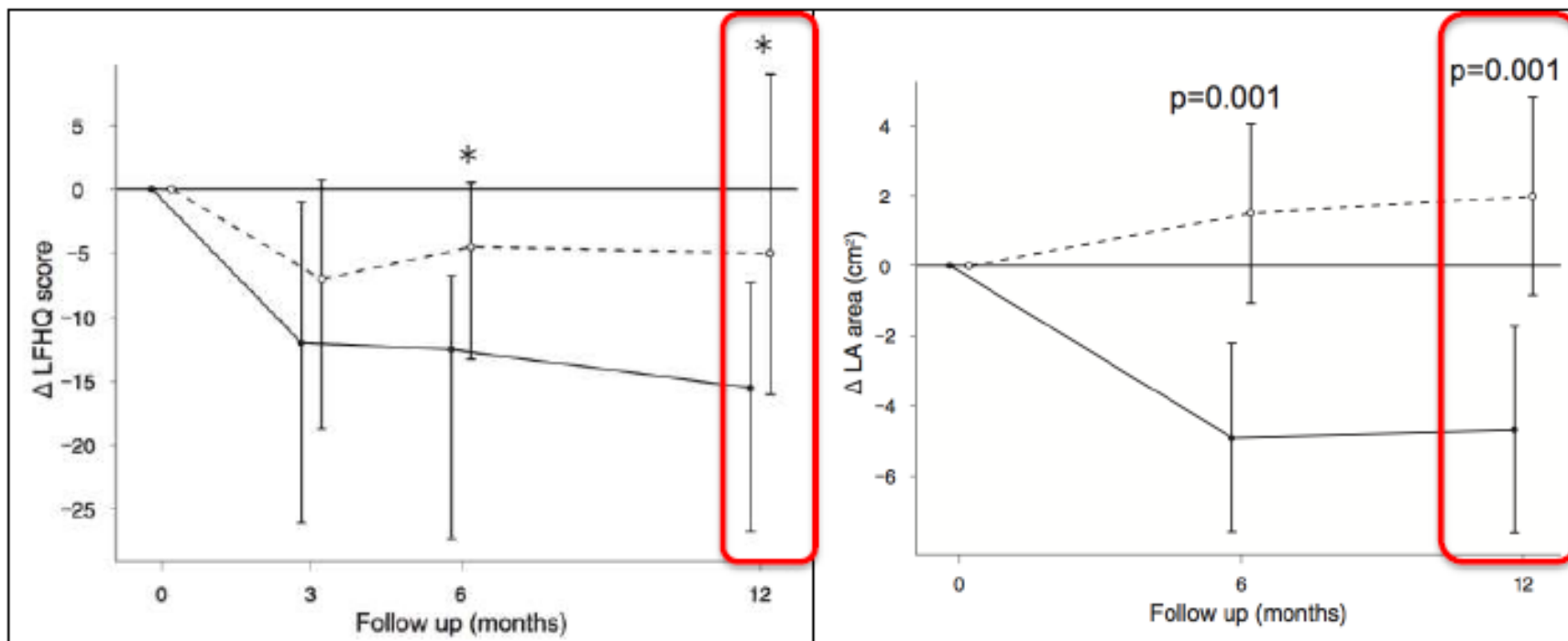
# Randomizovaná studie ARC-HF ablace perzist FiS x rate control

## Primary Endpoint



# Randomizovaná studie ARC-HF ablace perzist FiS x rate control

## Secondary Endpoints



# Randomizovaná studie ablace perzist FiS x AA medikace (CAMTAF)

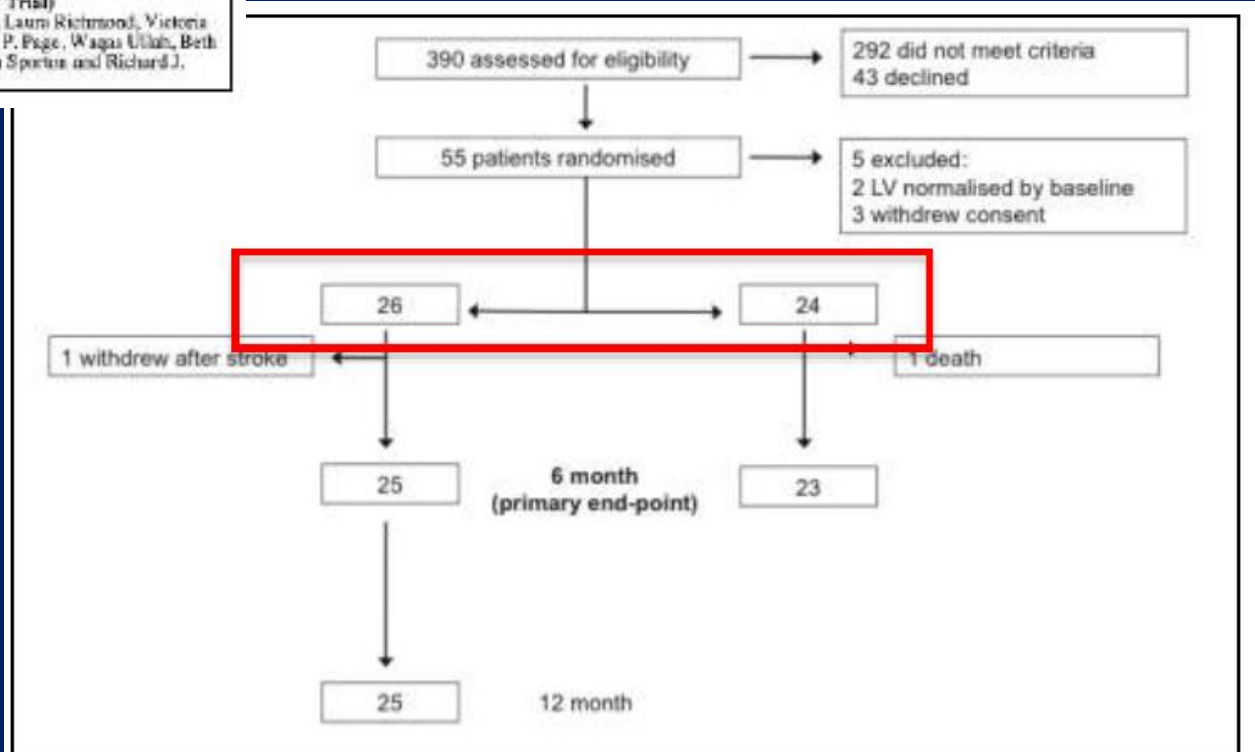
**Circulation**  
Arrhythmia and Electrophysiology



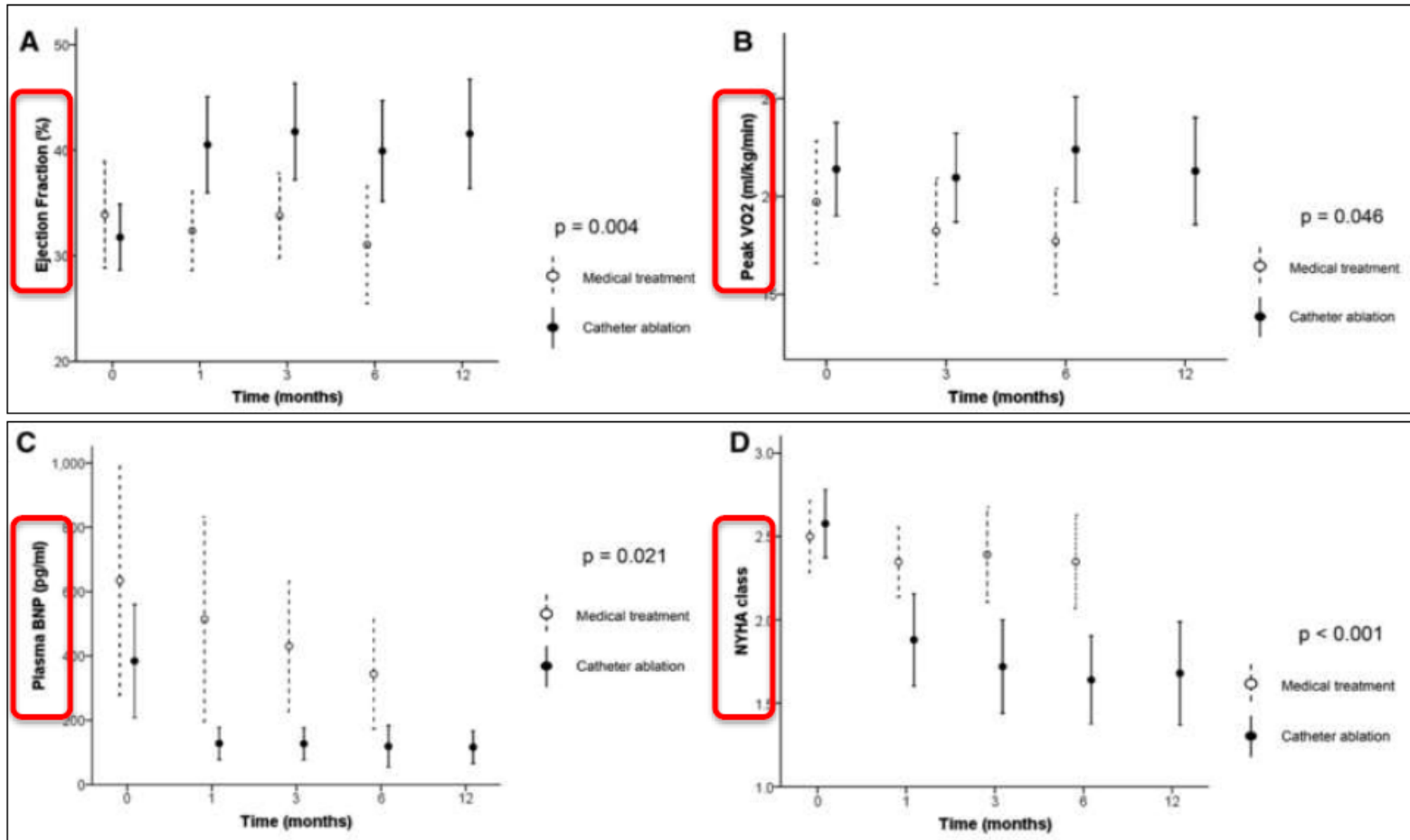
A Randomized Controlled Trial of Catheter Ablation Versus Medical Treatment of Atrial Fibrillation in Heart Failure (The CAMTAF Trial)  
Ross J. Hunter, Thomas J. Berman, Ihab Diab, Ravinolo Kannekar, Laura Richmond, Victoria Baker, Farai Gorenzoni, Vinit Sawhney, Edward Dawson, Stephen P. Page, Waqas Ullah, Beth Unsworth, Jamil Mujet, Mehul Chinoja, Mark J. Earley, Simon Spertus and Richard J. Schilling

EF pod 50% (prům 33%)  
Prim cíl : diff LVEF 6 Mo

Sek cíle:  
proBNP  
V02 max  
NYHA



# Randomizovaná studie ablace perzist FiS x AA medikace (CAMTAF)



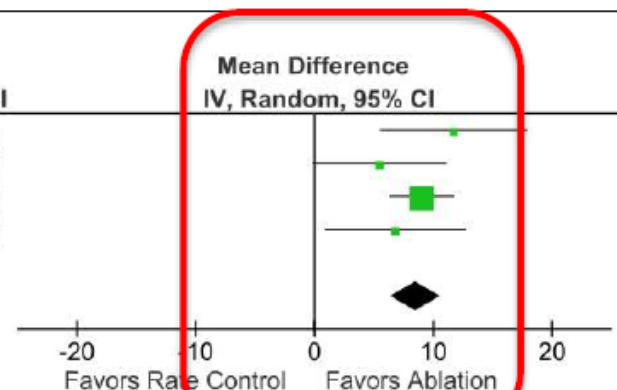
# Metaanalýza 4 randomizovaných studií perzist FiS x AA medikace u HF n= 224

## Change in left ventricular ejection fraction (LVEF)

**A**

Study or Subgroup	Ablation			Rate Control			Weight	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total		
Hunter 2014	8.1	12.5	26	-3.6	9.7	24	12.0%	11.70 [5.52, 17.88]
Jones 2013	10.9	11.5	24	5.4	8.5	26	14.3%	5.50 [-0.14, 11.14]
Khan 2008	8	8	41	-1	4	40	60.7%	9.00 [6.26, 11.74]
MacDonald 2011	8.2	12	20	1.4	5.9	18	13.0%	6.80 [0.88, 12.72]
<b>Total (95% CI)</b>			<b>111</b>			<b>108</b>	<b>100.0%</b>	<b>8.53 [6.40, 10.67]</b>

Heterogeneity: Tau<sup>2</sup> = 0.00; Chi<sup>2</sup> = 2.56, df = 3 (P = 0.46); I<sup>2</sup> = 0%  
 Test for overall effect: Z = 7.83 (P < 0.00001)

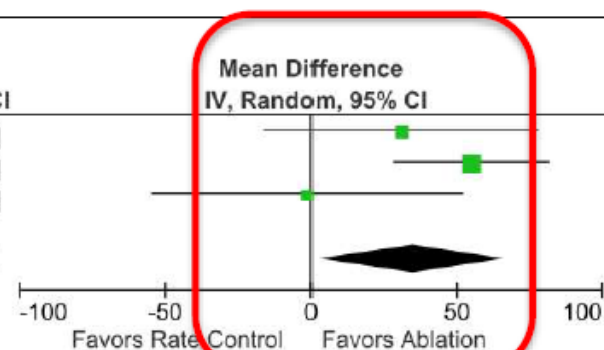


## Change in 6-min walk test distance

**C**

Study or Subgroup	Ablation			Control			Weight	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total		
Jones 2013	21	103.7	26	-10	65.19	26	28.2%	31.00 [-16.08, 78.08]
Khan 2008	71	78.48	41	16	40.2	40	47.9%	55.00 [27.94, 82.06]
MacDonald 2011	20.1	76.5	17	21.4	77.4	15	23.9%	-1.30 [-54.75, 52.15]
<b>Total (95% CI)</b>			<b>84</b>			<b>81</b>	<b>100.0%</b>	<b>34.76 [2.87, 66.65]</b>

Heterogeneity: Tau<sup>2</sup> = 362.12; Chi<sup>2</sup> = 3.61, df = 2 (P = 0.16); I<sup>2</sup> = 45%  
 Test for overall effect: Z = 2.14 (P = 0.03)



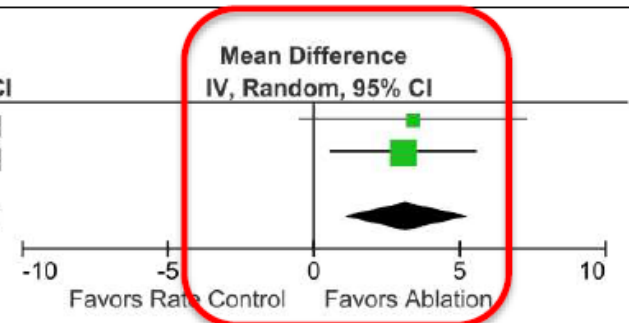
# Metaanalýza 4 randomizovaných studií perzist FiS x AA medikace u HF n= 224

## Change in peak oxygen consumption (VO2)

**D**

Study or Subgroup	Ablation			Rate Control			Weight	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total		
Hunter 2014	1.4	7.05	26	-2	7.1	24	29.1%	3.40 [-0.53, 7.33]
Jones 2013	2.13	5.52	24	-0.94	3.13	26	70.9%	3.07 [0.56, 5.58]
<b>Total (95% CI)</b>			<b>50</b>			<b>50</b>	<b>100.0%</b>	<b>3.17 [1.05, 5.28]</b>

Heterogeneity: Tau<sup>2</sup> = 0.00; Chi<sup>2</sup> = 0.02, df = 1 (P = 0.89); I<sup>2</sup> = 0%  
 Test for overall effect: Z = 2.93 (P = 0.003)

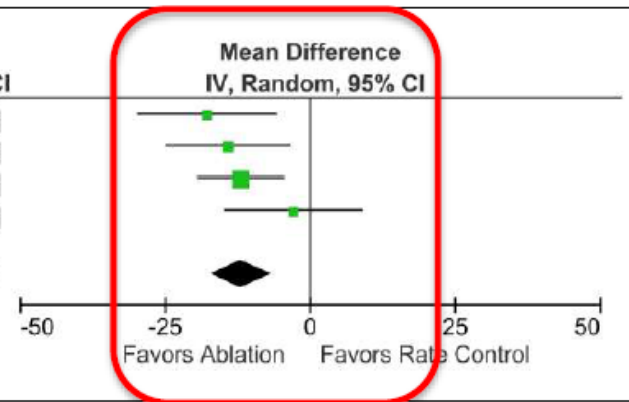


## Change in Minnesota Living With Heart Failure (MLWHF)

**B**

Study or Subgroup	Ablation			Rate Control			Weight	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total		
Hunter 2014	-18	21.99	26	-0.2	21.47	24	17.9%	-17.80 [-29.85, -5.75]
Jones 2013	-19.58	22.32	24	-5.35	15.71	26	22.0%	-14.23 [-25.01, -3.45]
Khan 2008	-19	21	41	-7	12.59	40	41.8%	-12.00 [-19.52, -4.48]
MacDonald 2011	-5.7	19.7	20	-2.8	17.9	18	18.2%	-2.90 [-14.85, 9.05]
<b>Total (95% CI)</b>			<b>111</b>			<b>108</b>	<b>100.0%</b>	<b>-11.88 [-17.15, -6.60]</b>

Heterogeneity: Tau<sup>2</sup> = 2.57; Chi<sup>2</sup> = 3.28, df = 3 (P = 0.35); I<sup>2</sup> = 8%  
 Test for overall effect: Z = 4.41 (P < 0.0001)





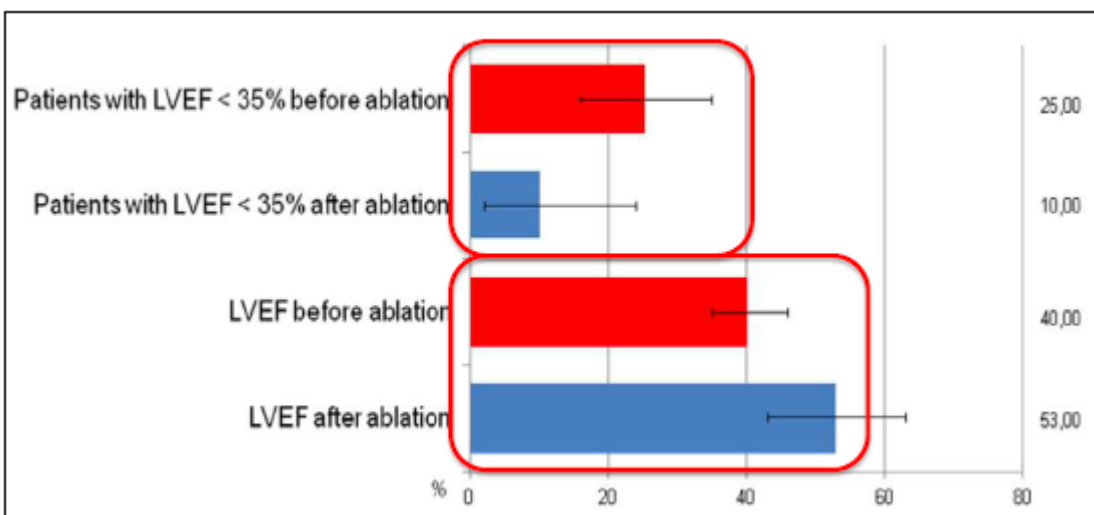
# Metaanalýza 23 studií perzist FiS x AA medikace u HF n= 1838, FUP 23měs

**Circulation**  
Arrhythmia and Electrophysiology



## Catheter Ablation of Atrial Fibrillation in Patients With Left Ventricular Systolic Dysfunction: A Systematic Review and Meta-Analysis

Matteo Anselmino, Mario Matta, Fabrizio D'Ascenzo, T. Jared Bunch, Richard J. Schilling, Ross J. Hunter, Carlo Pappone, Thomas Neumann, Georg Noelker, Martin Fiala, Emanuele Bertaglia, Antonio Frontera, Edward Duncan, Chishan Nalliah, Pierre Jais, Ruksben Weerasooriya, Jon M. Kalman and Fiorenzo Gaita



# Mortalitní randomizovaná studie Ablace Perzist FiS x amiodarone (AATAC)

- Multicent, 2D-ICD, CRT-D, EF pod 40%,
- RMT 102:102 pts

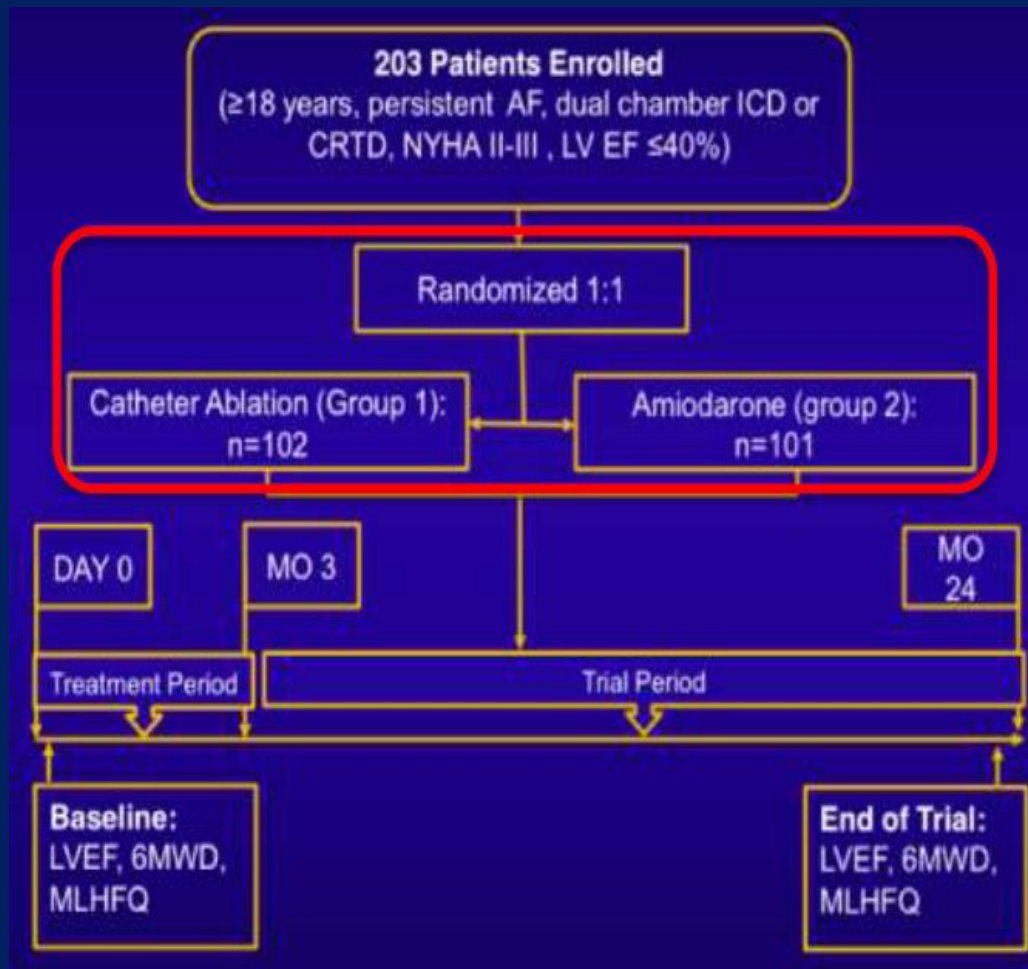
Prim cíl: AF rekurence

Sek cíl: mortalita,

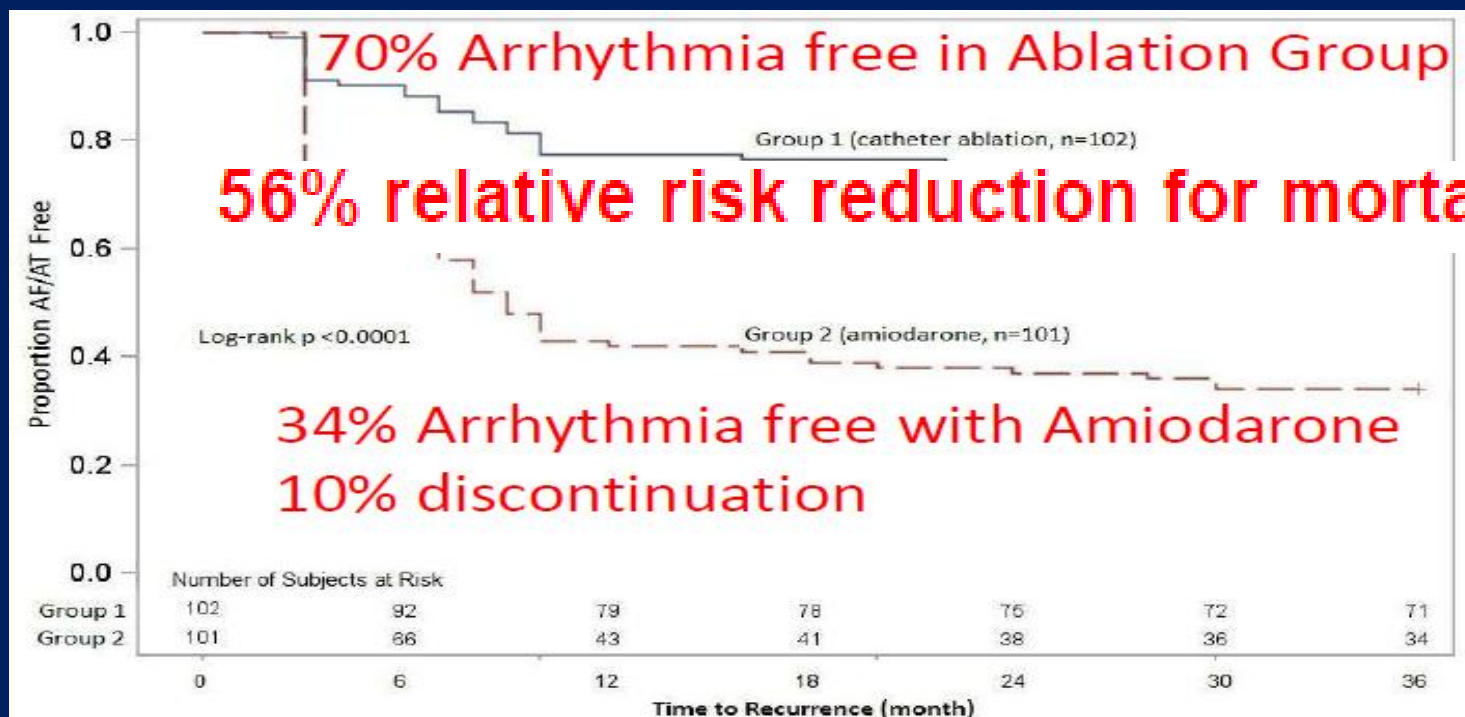
Hospitalizace, 6minWD

MLHFQ

FUP 24měs



# Mortalitní randomizovaná studie Ablace Perzist FiS x amiodarone (AATAC)



	No Recurrence (n= 91)		Recurrence (n= 86)		P (comparing change between groups)
	Baseline	Change (median)	Baseline	Change (median)	
LVEF (%)	28.8±10	9.6±7.4 (9.4)	30.2±9	4.2±6.2 (4.0)	<0.001
6MWD (meter)	347±113	27±38 (24)	352±128	8±42 (2)	<0.001
MLHFQ	53±24	-14±18 (-12)	49±26	-2.9±15 (-2.2)	<0.001

# 6-leté výsledky ablace FiS EF<45%

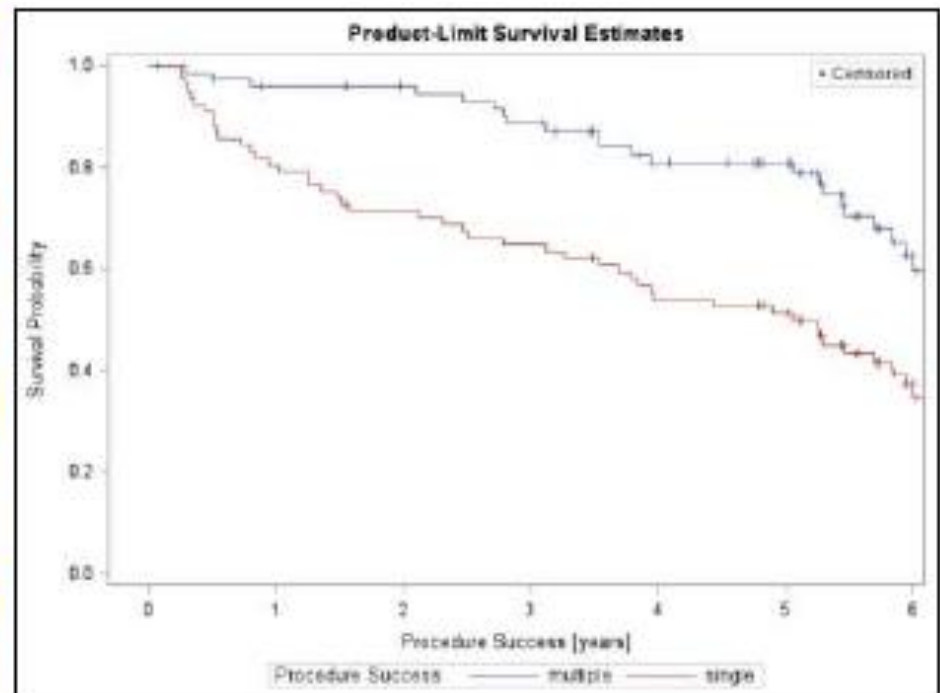
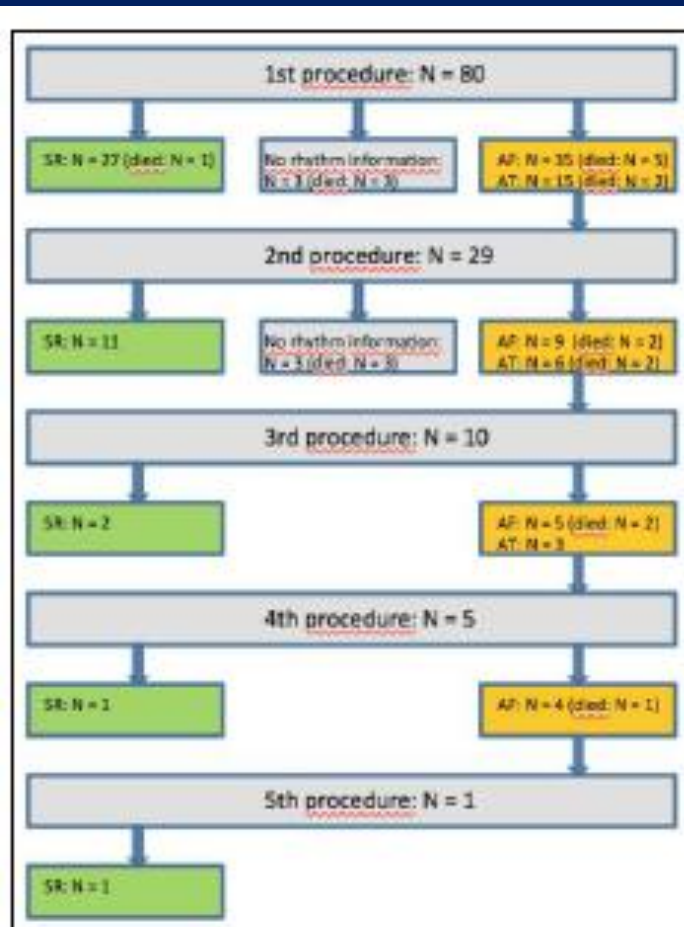
## Six-Year Clinical Outcomes After Catheter Ablation of Atrial Fibrillation in Patients With Impaired Left Ventricular Function

ANDREAS RILLIG, M.D.,\* HISAKI MAXIMOTO, M.D.,\* JASCHA WEGNER,\* TINA LIN, M.B.B.S., B.MED.SCI., F.R.A.C.P.,\* CHRISTIAN HEEGER, M.D.,\* CHRISTINE LEMES, M.D.,\* THOMAS FINK, M.D.,\* ANDREAS METZNER, M.D.,\* ERIK WISSNER, M.D.,\* SHIBU MATHREW, M.D.,\* PETER WOILMUTH† KARL-HEINZ RUCK, M.D.,\* ROLAND RICHARD TILZ, M.D.,\* and FEIFAN OUYANG, M.D.\*

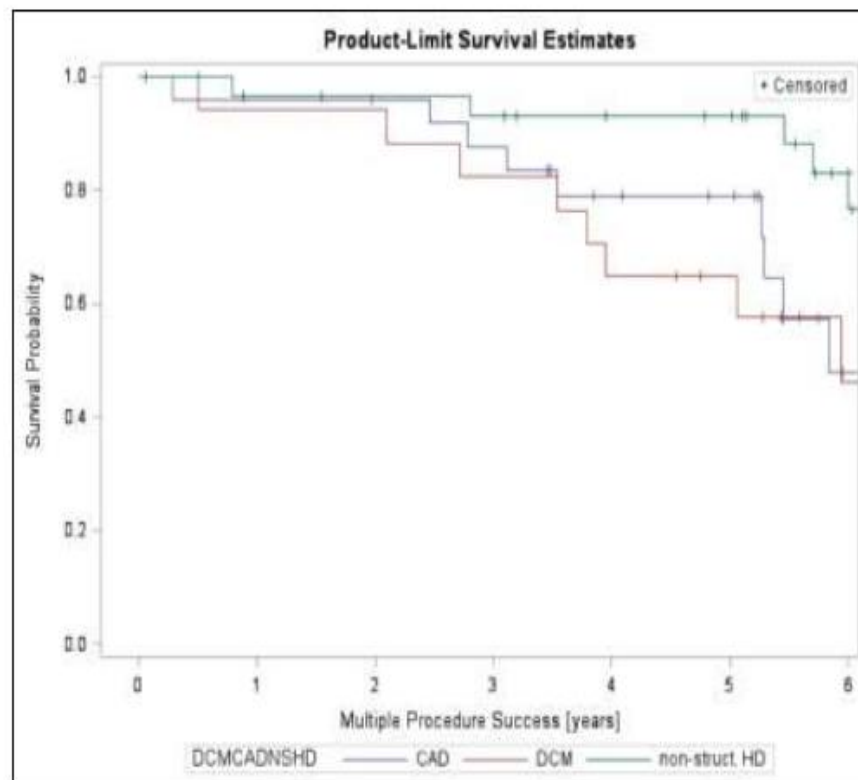
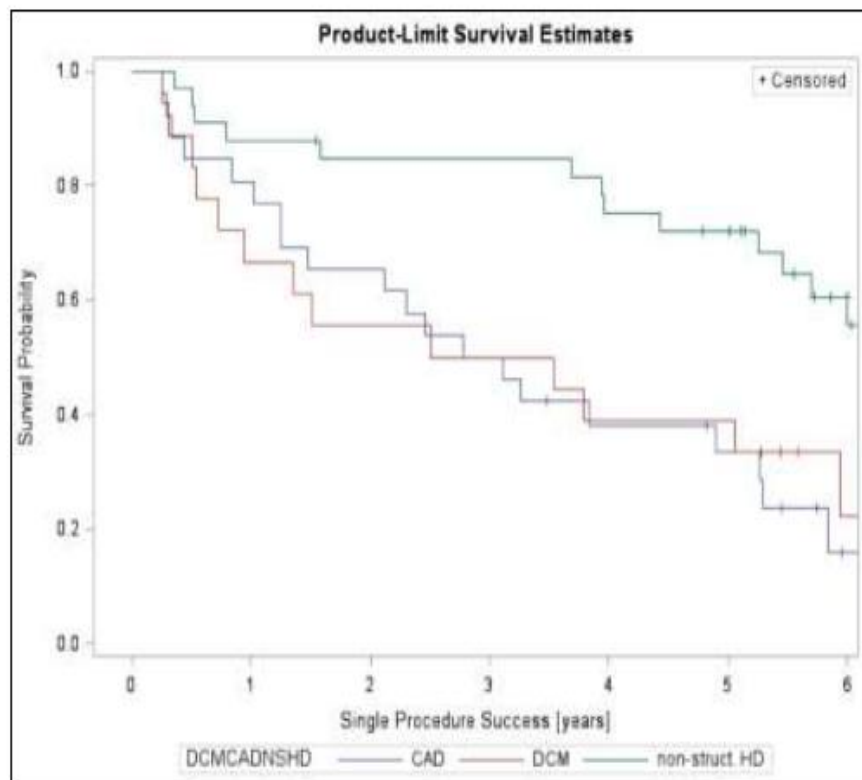
From the \*Department of Cardiology, Asklepios Klinik St. Georg, Hamburg, Germany; and †Asklepios Proresearch, Hamburg, Germany

- N= 80 ,
- Parox 20%, perzist 46%, dlouhod. perzist 33%
- ICHS, DKMP, tachyKMP
- FUP median 72m (67-75), Holter EKG, KS/ICD

# 6-leté výsledky ablace FiS EF < 45%

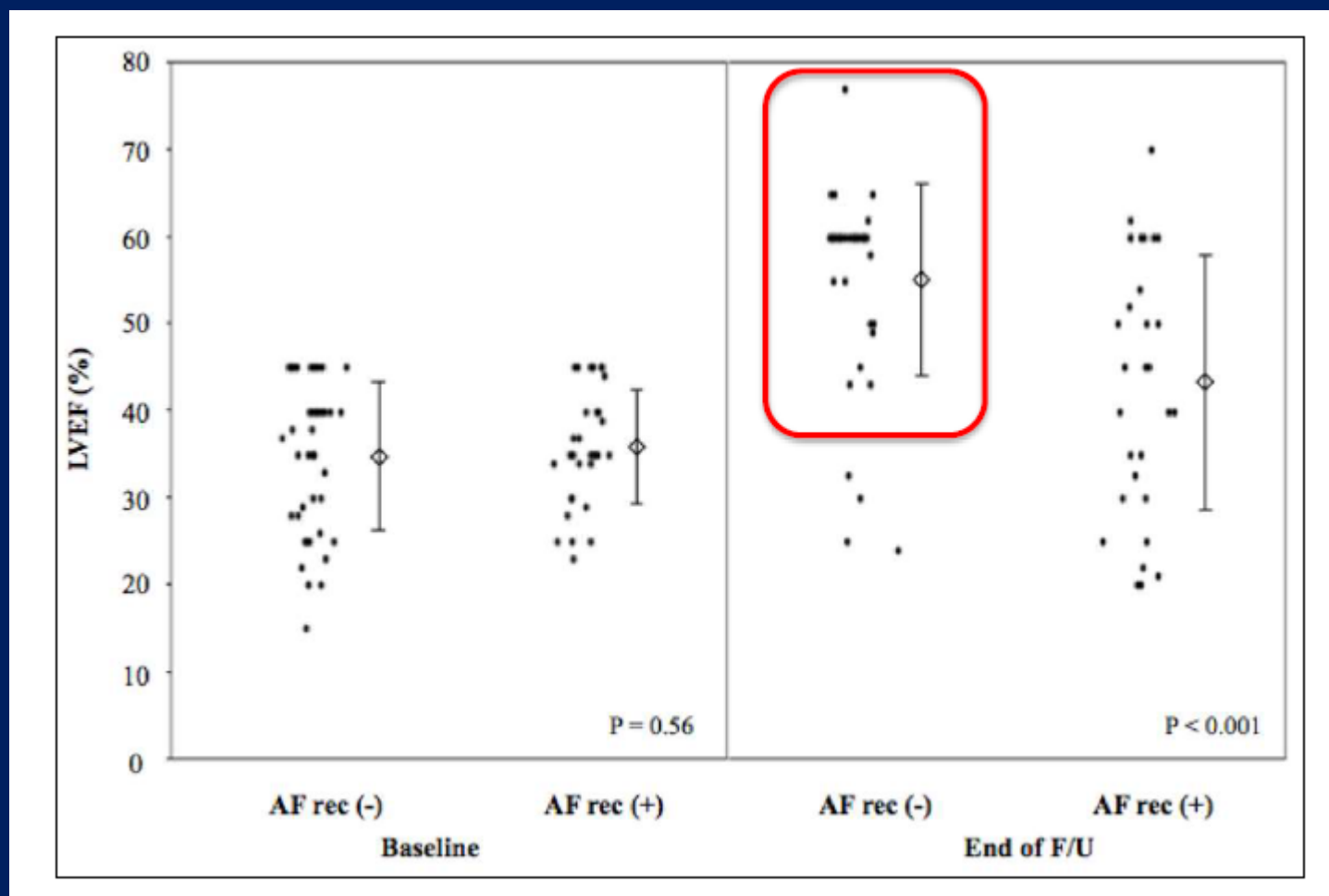


# 6-leté výsledky ablace FiS EF<45%



# 6-leté výsledky ablace FiS EF < 45%

Statisticky významné zlepšení EFLK,  $p < 0.001$





# 5 letá mortalita a CMP po první ablací FiS u srdečního selhání

- Srovnání 3 skupin 1:4:4 s věkovou korelací
- 1. HF-FiS první RFA n=267
- 2. HF-FiS bez RFA n=1068
- 3. HF bez FiS n=1068
- EF < 35%
- Prim cíl: celk.mortalita, CMP, EFLK-NYHA, recid FiS

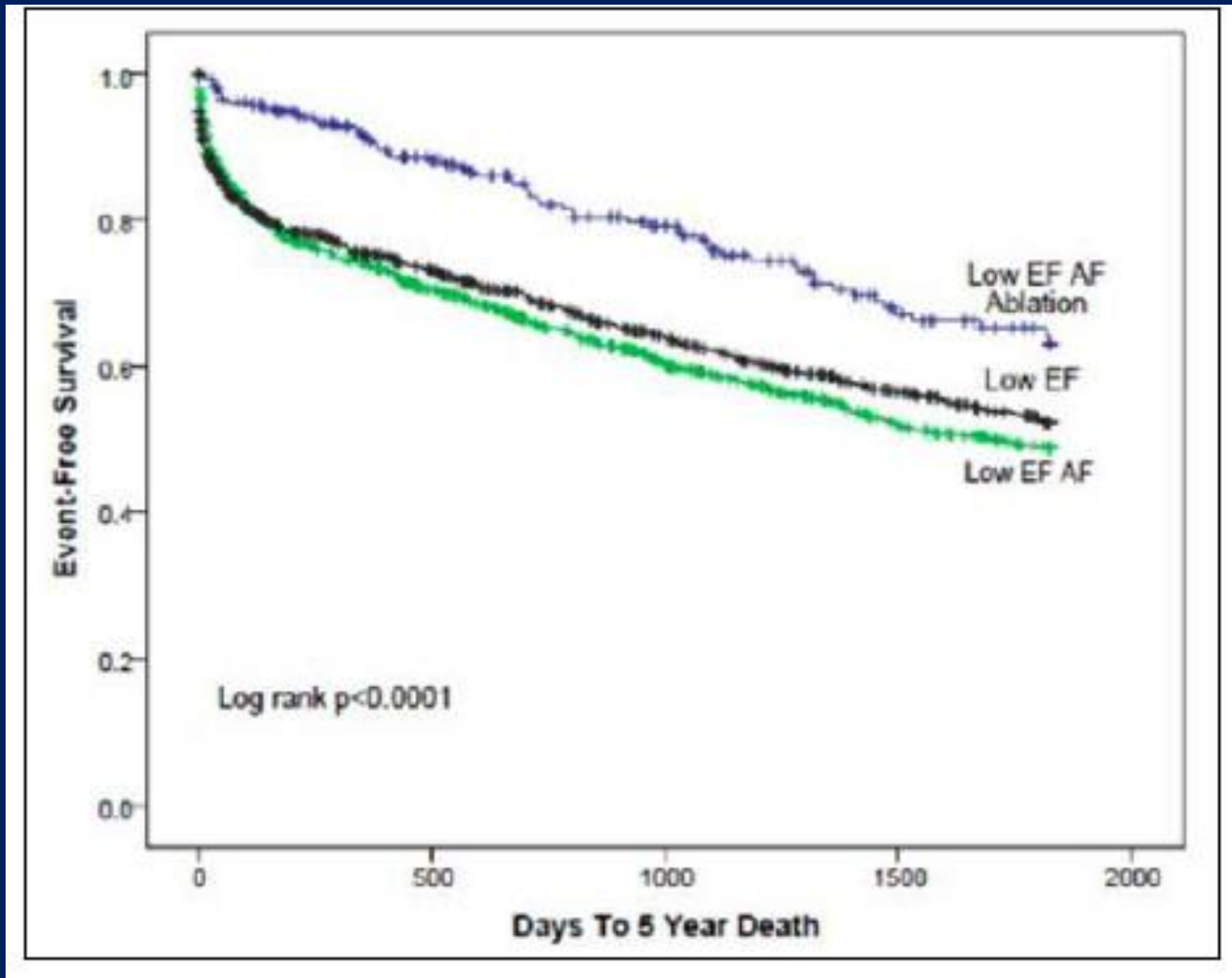
## Five-Year Outcomes of Catheter Ablation in Patients with Atrial Fibrillation and Left Ventricular Systolic Dysfunction

T. JARED BUNCH, M.D., HEIDI T. MAY, Ph.D., M.S.P.H., TAMI L. BAIR, R.N., VICTORIA JACOBS, N.P., BRIAN G. CRANDALL, M.D., MICHAEL CUTLER, D.O., Ph.D., J. PETER WEISS, M.D., CHARLES MALLENDER, M.D., JEFFREY S. OSBORN, M.D., JEFFREY L. ANDERSON, M.D., and JOHN D. DAY, M.D.

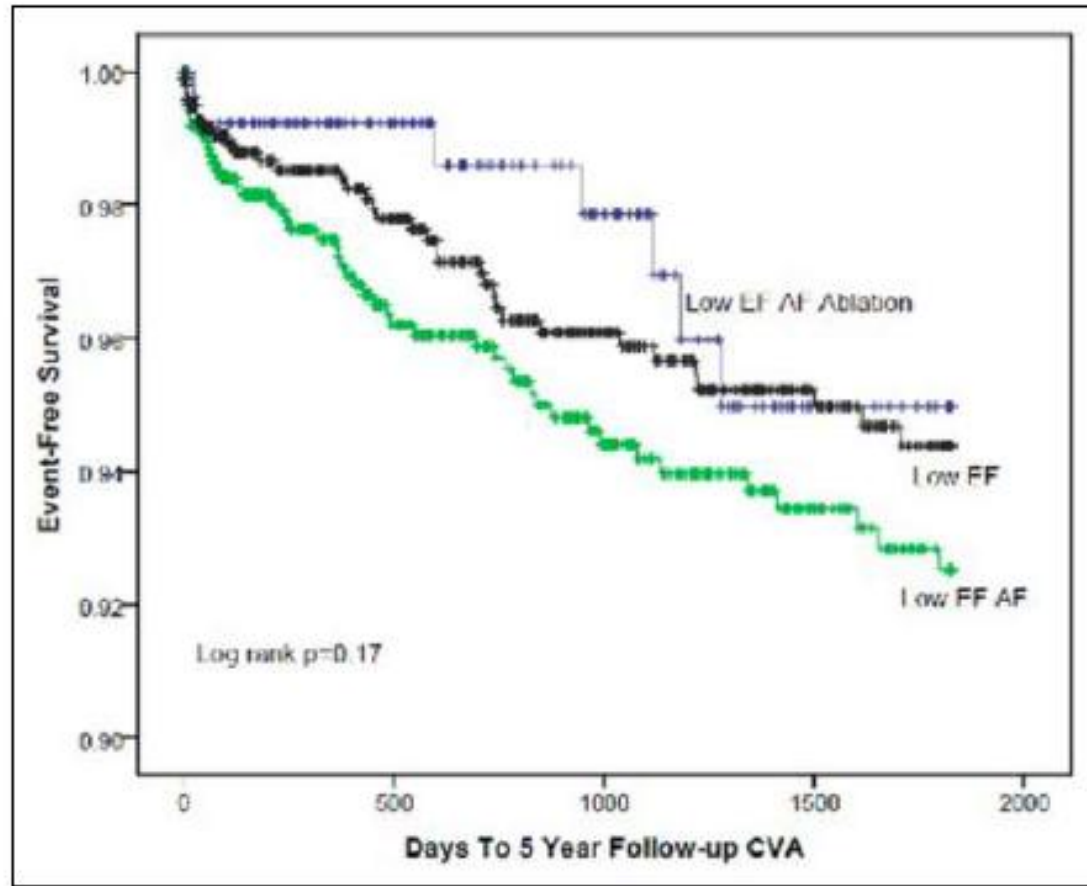
From the Intermountain Heart Institute, Intermountain Medical Center, Murray, Utah, USA



# 5 letá mortalita po první ablaci FiS u srdečního selhání

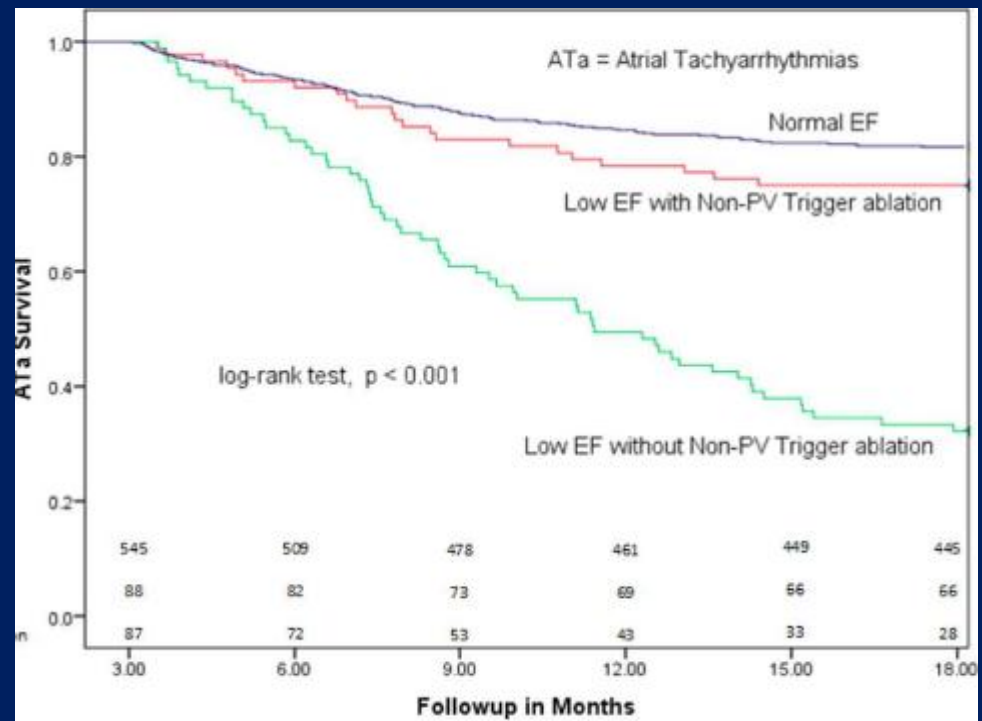
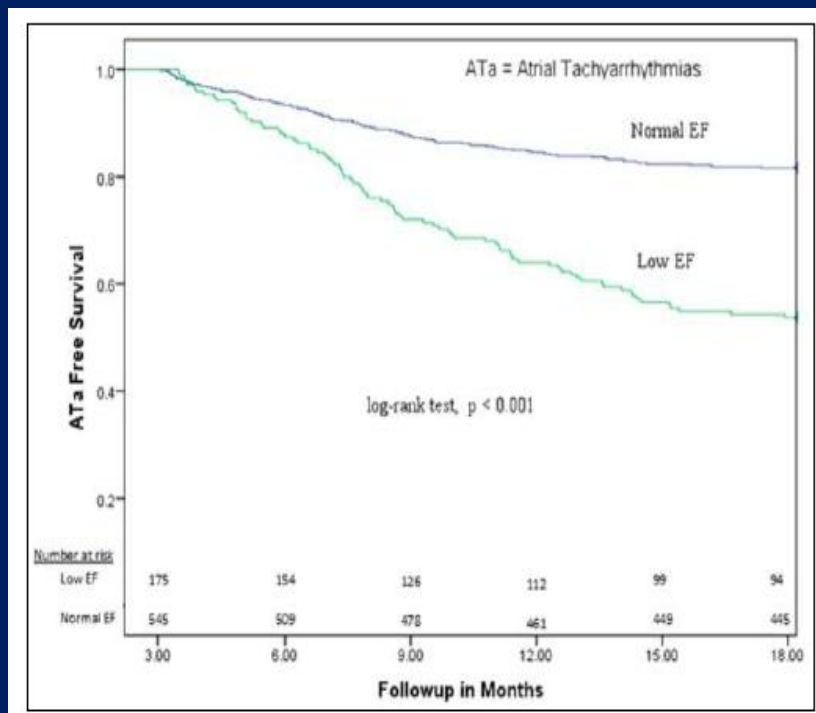


# 5 letý výskyt CMP po ablaci FiS u srdečního selhání

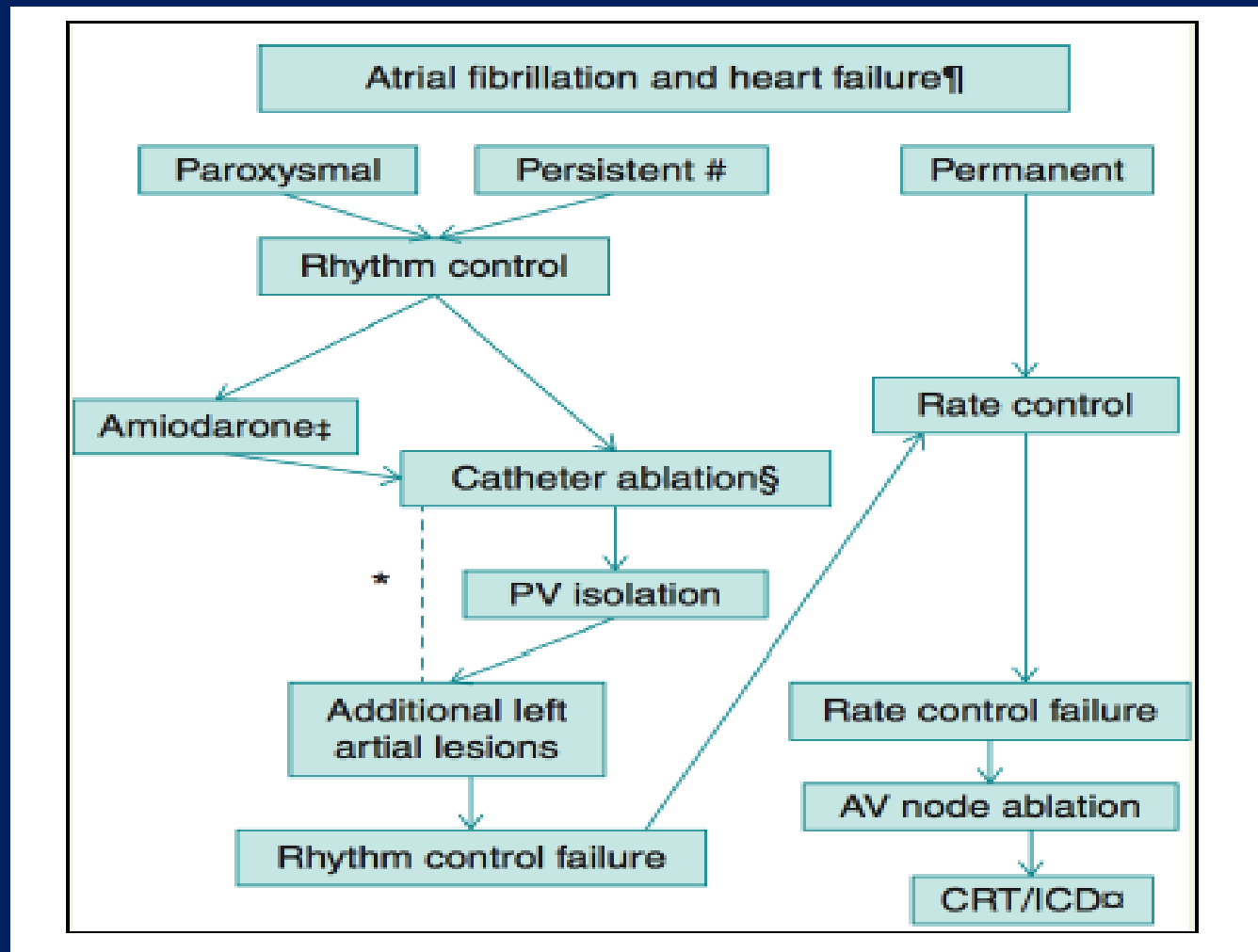


# Jak zlepšit úspěšnost ablace FiS u srdečního selhání ?

## Ablace extrapulmonálních spouštěčů FiS



# Algoritmus co s FiS u srd. selhání



# Závěr

- Cílenou katetrizační ablací fibrilace síní u srdečního selhání lze očekávat významné zlepšení kvality života, fyzické výkonnosti, redukce CMP, významný nárůst EF LK a významný mortalitní benefit.
- Optimálních výsledků lze dosáhnout komplexním – individuálním přístupem (extrapulmonální spouštěče, terminace FiS ablací, noninducibilita).
- Rozvaha o implantaci CRT-D případně s neselektivní ablací AV junkce by měla být zvažována až ve druhé době po vyhodnocení ev. neúspěchu primární strategie

# **Role katetrizační ablace fibrilace síní v léčbě srdečního selhání**

**Jan Škoda**

**Nemocnice Na Homolce**

