



IV Congresso
Novas Fronteiras
em Cardiologia

Cardiopatia estrutural. O que há de novo?

Encerramentos de CIA, FOP e AAE

7 a 9 de Fevereiro 2014
Hotel Vila Galé Ericeira

Marco Costa
CHUC-HG



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Novas Fronteiras em Cardiologia

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Cardiopatia estrutural. O que há de novo?

1. *Encerramentos de CIAOs*

- *Erosão cardíaca peridispositivo*

2. *Encerramentos de FOPs*

- *Estudo RESPECT*

3. *Encerramentos de AAE*

- *Registo clínico*

Marco Costa
CHUC-HG

CORRECÇÃO DE DEFEITOS DO SEPTO INTERAURICULAR

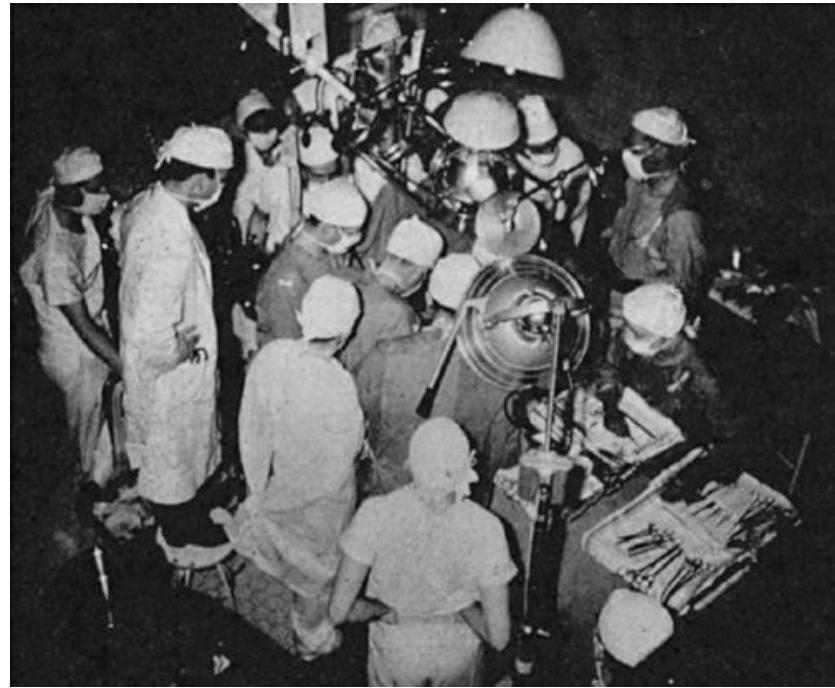
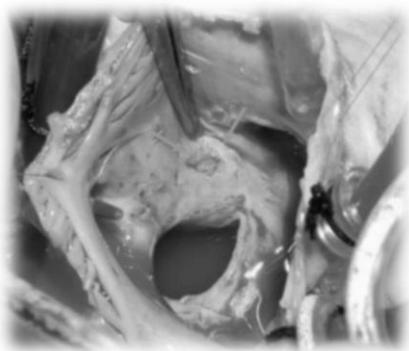
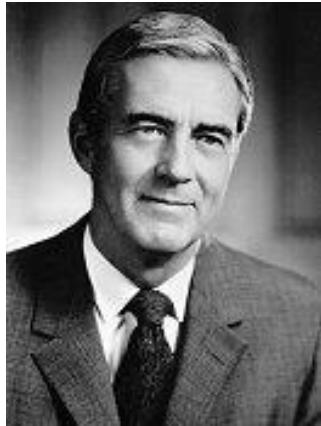
Perspectiva história



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1950

Encerramento cirúrgico de CIA





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CORRECÇÃO DE DEFEITOS DO SEPTO INTERAURICULAR

Perspectiva história

1950

Encerramento cirúrgico de CIA

1974

Encerramento percutâneo de CIA



Secundum Atrial Septal

Nonoperative Closure During Cardiac Catheterization

Terry D. King, MD; Sandra L. Thompson, RN; Charles

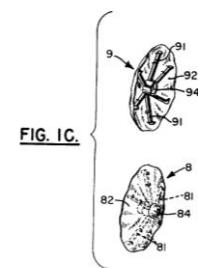
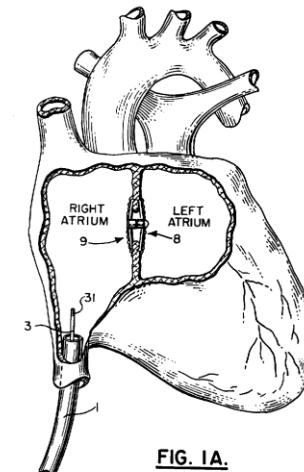
- A 17-year-old girl had clinical and cardiac catheterization compatible with a secundum atrial septal defect. During cardiac catheterization the atrial septal defect was sized and closed using a transseptal technique.

(JAMA 235:2506-2509, 1976)

PATENTED APR 1 1975

3,874,388

SHEET 01 OF 10





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CORRECÇÃO DE DEFEITOS DO SEPTO INTERAURICULAR

Perspectiva história

1950

Encerramento cirúrgico de CIA

1974

Encerramento percutaneo de CIA

1992

Encerramento de FOP após AVC criptogénico

Circulation

Transcatheter closure of patent foramen ovale after presumed paradoxical embolism.

N D Brdaes; W Hellenbrand; L Latson; J Filiano; J W Newburger; J E Lock

1992;86:1902-8. 10.1161/01.CIR.86.6.1902

Abstract

BACKGROUND Many have proposed a relation between presence of a patent foramen ovale, with or without atrial septal aneurysm, and cryptogenic stroke. The effect of foramen ovale closure on the risk for subsequent strokes is unknown.

METHODS AND RESULTS Transcatheter closure of a patent foramen ovale was undertaken in 36 patients with known right-to-left atrial shunting and presumed paradoxical emboli (31 strokes, 25 transient neurological events, four systemic arterial emboli, and two brain abscesses). Individual patients had between one and four such events. None had a left heart or carotid source of embolism; 31 of 35 had no known risk factors for stroke. Events occurred in 12 patients while they were taking warfarin. At cardiac catheterization, patent foramina ovalia were significantly larger than predicted for age in 67% of the patients. Implantation of a double-umbrella device in the patent foramen ovale was achieved in all without serious procedural complications. Of 34 who have returned for follow-up, one has a residual atrial communication that may be clinically important, five had trivial leaks, and 28 have complete closure. There have been no strokes during a mean follow-up of 8.4 months.

CONCLUSIONS Transcatheter closure of a patent foramen ovale can be accomplished with little morbidity and may reduce the risk of recurrence. Further investigations directed toward identifying the population at risk and assessing the effect of intervention are warranted.

Copyright © 1992 by American Heart Association





INDICAÇÕES PARA ENCERRAMENTO DE COMUNICAÇÃO INTERAURICULAR

4.1 Atrial septal defect

Introduction and background

ASD may not uncommonly remain undiagnosed. ASD types include:

- B Secundum ASD (80% of ASDs; located in the ovalis and its surrounding)
- D Primum ASD [15%, synonyms: partial atrial defect (AVSD), partial atrioventricular canal], at the crux, AV valves are typically normal in various degrees of regurgitation; see Secundum ASD
- A Superior sinus venosus defect [5%, located near vena cava (SVC) entry, associated with connection of right pulmonary veins to SVC]
- C Inferior sinus venosus defect [<1%, located near vena cava (IVC) entry]
- E Unroofed coronary sinus [<1%, separate from LA] can be partially or completely missing

ESC Guidelines for the management of grown-up congenital heart disease (new version 2010)

Table 3

Indications for intervention in atrial septal defect

Indications	Class ^a	Level ^b
Patients with significant shunt (signs of RV volume overload) and PVR <5 WU should undergo ASD closure regardless of symptoms	I	B ^{c,d}
Device closure is the method of choice for secundum ASD closure when applicable	I	C
All ASDs regardless of size in patients with suspicion of paradoxical embolism (exclusion of other causes) should be considered for intervention	IIa	C
Patients with PVR ≥5 WU but <2/3 SVR or PAP <2/3 systemic pressure (baseline or when challenged with vasodilators, preferably nitric oxide, or after targeted PAH therapy) and evidence of net L–R shunt ($Qp:Qs > 1.5$) may be considered for intervention	IIb	C
ASD closure must be avoided in patients with Eisenmenger physiology	III	C



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Benefits of Transcatheter Closure

- No exposure to cardiopulmonary bypass
- No chest incision required
- Decreased need for blood or blood product transfusion
- Reduction in hospital stay
- Significantly reduced convalescence time
- Rapid return to normal activities
- Potential health care economic benefits

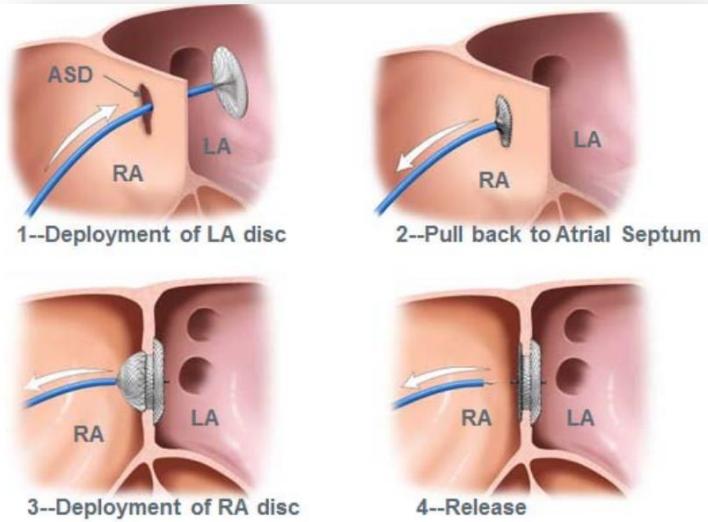


Table 1. All-Cause Mortality by Treatment Modality

All-Cause Mortality			
Reference	Surgical Closure	Transcatheter Closure	AMPLATZER ASO Transcatheter Closure
Karamlou ⁴	0.88%	0.60%	0.009%*
DiBardino ⁵	0.13%	0.093%	

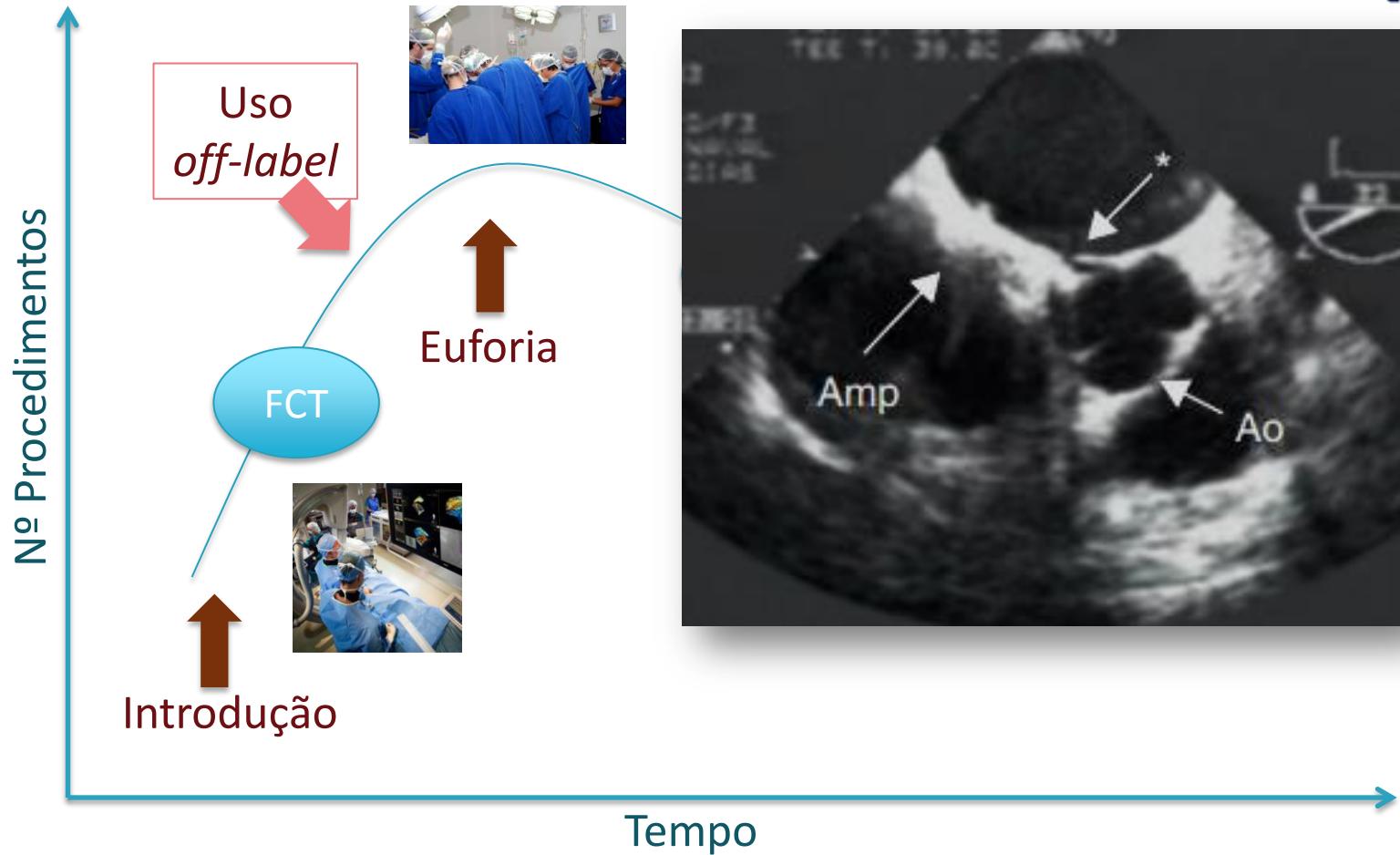
*0.009% is based on post commercial experience. Reference Section 4.

Atrial Septal Occluder Executive Summary
www.fda.gov/downloads/.../UCM304944.pdf
24/05/2012

NOVAS TECNOLOGIAS E DISPOSITIVOS NA CARDIOLOGIA DE INTERVENÇÃO



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ST. JUDE MEDICAL
MÉDICO CONTROLE DIRETO

St. Jude Medical
AMPLATZER® Atrial Septal Occluder
Executive Summary

Advisory Committee Briefing Materials: Available for Public Release

St. Jude Medical
24 May 2012

World-Wide Serious Adverse Event Overview – Through January 2012*

Adverse Event	Total Reported Events (on-label, ASO, world-wide)	Rate (based on sales, world-wide)**
Arrhythmia	54	0.024%
Embolization	347	0.155%
Erosion	97	0.043%
Fracture	1	0.0004%
Malfunction	0	0.000%
Malposition	9	0.004%
Stroke	6	0.003%
Thrombus on Device	11	0.005%

*Erosion events are as of 15 March 2012

** All adverse event rates are calculated using total world-wide sales of 223,965 units.

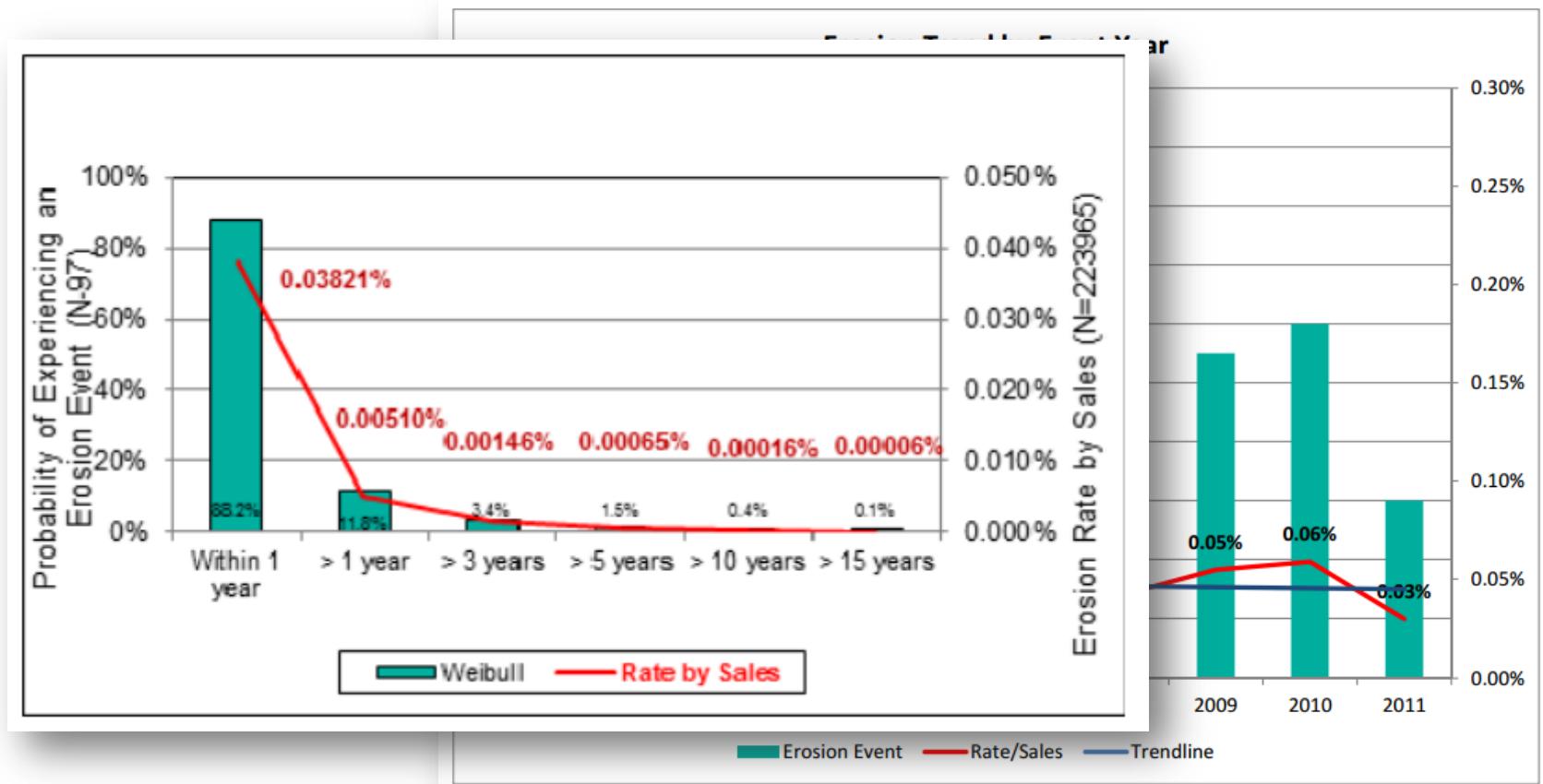
NOTE: The same patient may be counted in more than one of the above reported adverse events.

Atrial Septal Occluder Executive Summary
www.fda.gov/downloads/.../UCM304944.pdf 24/05/2012



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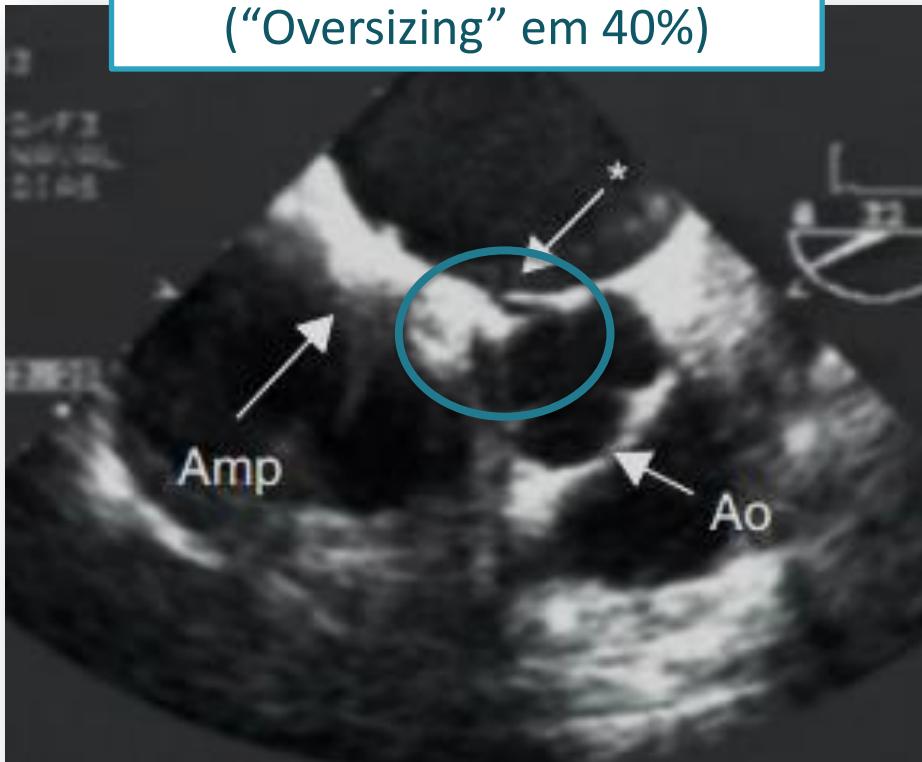
Atrial Septal Occluder Executive Summary
www.fda.gov/downloads/.../UCM304944.pdf 24/05/2012



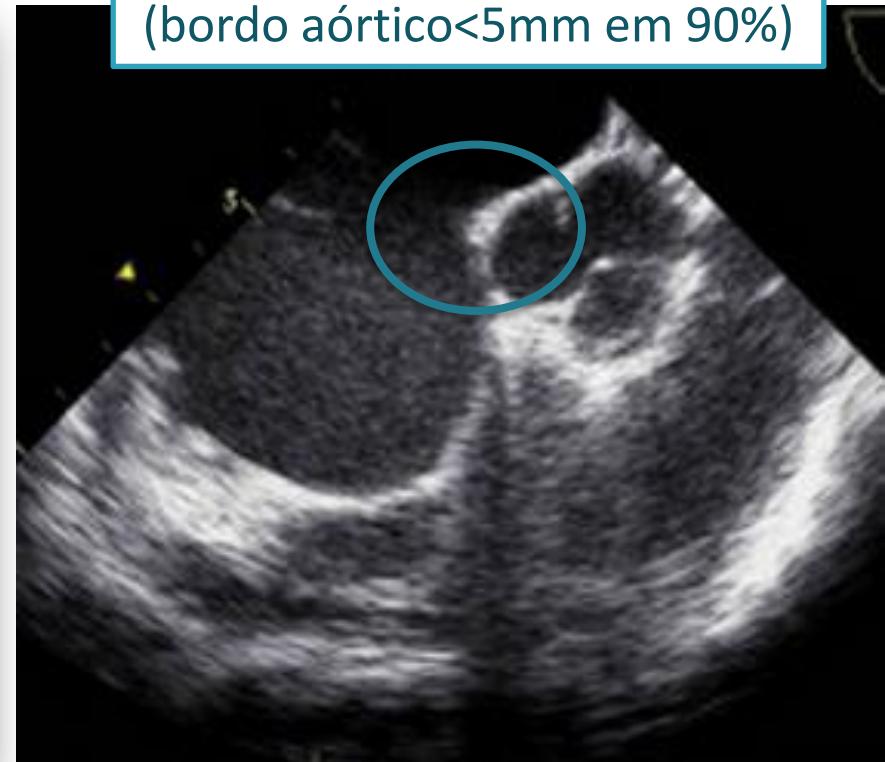
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CAUSAS DE EROSÃO APÓS ENCERRAMENTO DE CIA

TAMANHO INAPROPRIADO
("Oversizing" em 40%)



BORDOS INSUFICIENTES
(bordo aórtico<5mm em 90%)





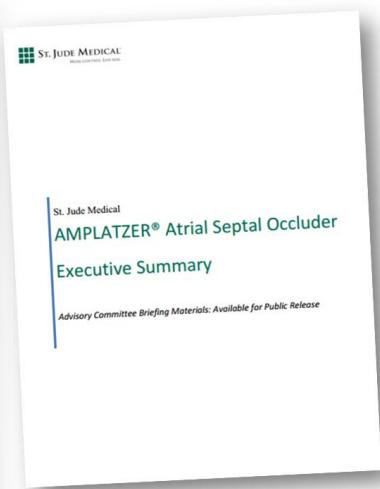
O QUE HÁ DE NOVO NO ENCERRAMENTO DE COMUNICAÇÃO INTERAURICULAR

Additional contraindications:

- Any patient in whom the device would interfere with or contact other intracardiac or intravascular structures (e.g. atrial roof, cardiac valves, pulmonary veins, coronary sinus, or aorta).
- Any patient with echocardiographic evidence of absent or deficient anterior – superior rim (sufficient rim is defined as the presence of at least 5mm of rim in multiple AND sequential short axis views confirmed by ICE or TEE).

Updated warning:

Do not release the AMPLATZER Septal Occluder from the delivery cable if the device does not conform to its original configuration, or if the device position is unstable and/or in contact with any adjacent cardiac structure.



Atrial Septal Occluder Executive Summary
www.fda.gov/downloads/.../UCM304944.pdf 24/05/2012

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(...)The ACC believes that there is insufficient data and no consensus as to whether or how to change clinical practice or alter labeling of ASD occlusion devices.

(...) In reviewing the literature and available registry data, it is evident that there is neither conclusive data nor consensus about incidence or root cause(s) of cardiac perforation or erosion by the ASO. Potential risk factors are:

- Rotation of the device around its central pins during atrial contraction (translational movement of the device relative to the motion of the heart) after implantation
- Splaying or flaring of the device around the aortic root following implantation
- Contact by the edge of the device with the atrial wall causing protrusion of the device into wall and into adjacent structures such as the aorta
- Absent and/or deficient aortic (anterior-superior) rim
- Thicker device profile at the time of deployment.

These factors individually or in combination may be predictors of early and late erosions; thus, they warrant close monitoring.



O QUE HÁ DE NOVO NO ENCERRAMENTO DE COMUNICAÇÃO INTERAURICULAR



Informação Importante sobre Dispositivo Actualização das Instruções de Utilização (IFU) com o Oclusor Septal AMPLATZER™

22 de Novembro de 2013

Exmo. Sr. Dr Marco Costa

Vimos pela presente transmitir-lhe informação importante sobre actualização das instruções de utilização do Oclusor Septal AMPLATZER (ASO). Após muitos anos de experiência com o St. Jude Medical vem agora actualizar as instruções relativas ao ASO, de modo a refletir a experiência clínica mais recentes. Estas alterações destinam-se a fornecer orientações para a utilização do dispositivo ASO e para atenuar a ocorrência de casos de erosão grave. A análise de eventos de erosão, confirmados ou potenciais, sobre contra-indicações e avisos, e fornecer também orientações adicionais e recomendações de seguimento que são apresentadas nas IFU. O objectivo é fornecer-lhe informações sobre essas alterações.

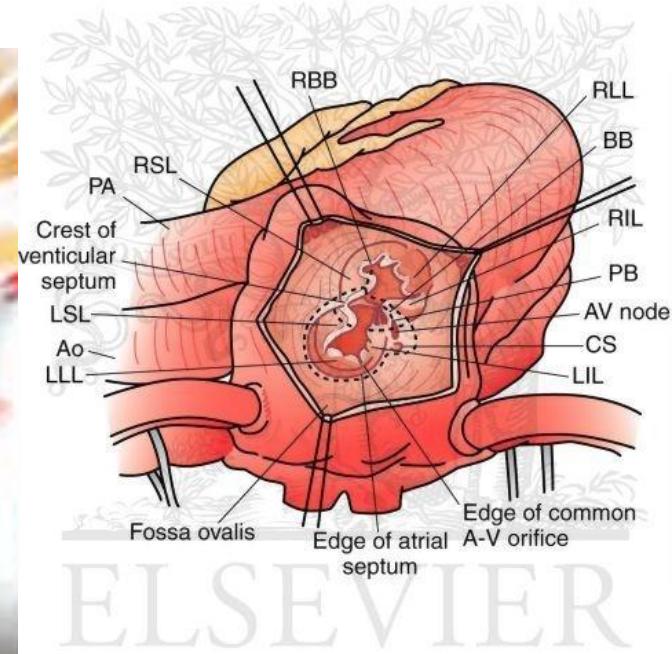
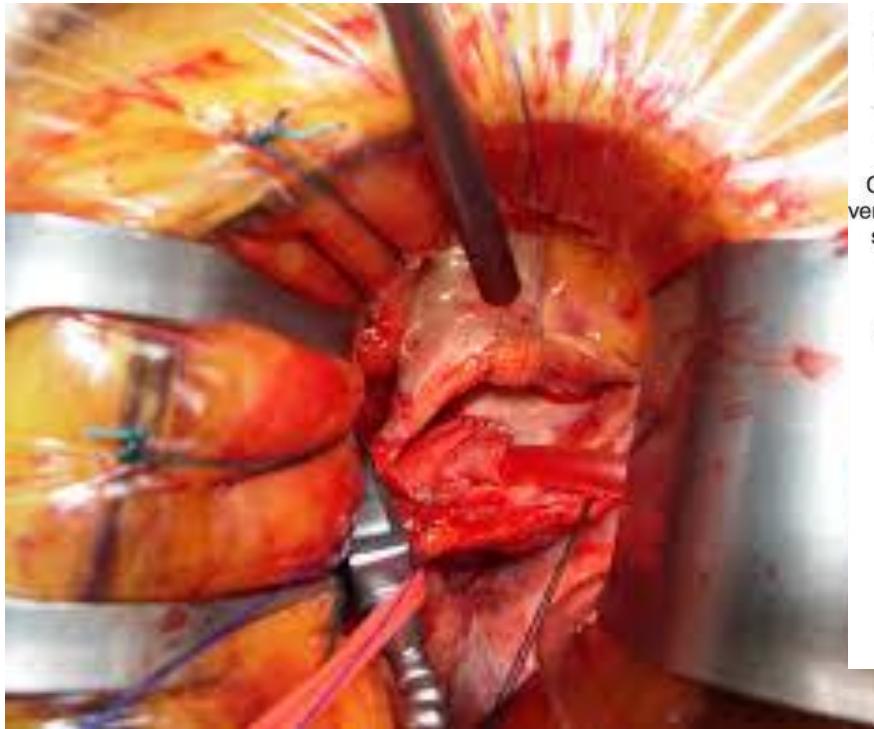
Produtos afectados	Oclusor Septal AMPLATZER™ (ASO) Modelos # ASD-004 a 040
Evento Adverso	A erosão, apesar de rara, é um evento potencialmente grave, com risco de vida, cuja sintomatologia inclui dor torácica, falta de ar, desmaios e dificuldades respiratórias. Caso se verifique uma erosão, poderá ser necessária uma cirurgia de emergência para assegurar um resultado positivo.
Especificamente erosão	
Risco de erosão	O risco de potencial erosão não tem sofrido alterações ao longo do tempo. A taxa de incidência global dos casos de erosão situa-se entre os 0,1% e os 0,3%.
Revisão das Instruções de Utilização do ASO	As principais revisões das IFU acrescentam informação adicional para os médicos, relativamente os riscos e sintomas da erosão. As IFU receberam as seguintes actualizações e esclarecimentos: <ul style="list-style-type: none">• A contra-indicação sobre margens de defeito inferiores a 5 mm foi actualizada, de modo a incluir o bordo da veia cava inferior• Os avisos foram actualizados ou modificados para incluir o seguinte:<ul style="list-style-type: none">○ Os doentes com um bordo retro-aórtico inferior a 5 mm em qualquer plano ecocardiográfico ou os doentes nos quais o dispositivo atinge a raiz da aorta podem estar sujeitos a um maior risco de erosão○ A colocação do oclusor ASD pode afectar intervenções cardíacas futuras, por exemplo, uma punção transseptal e a reparação da válvula mitral○ Não soltar o dispositivo do cabo de entrega se o dispositivo não estiver conforme a respectiva configuração original, ou se o dispositivo não estiver numa posição estável ou se interferir com qualquer estrutura cardíaca adjacente, como a veia cava superior (VCS), válvula pulmonar (VP), válvula mitral (MV), seio coronário (SC) ou aorta (AO). Recapturar o dispositivo e implantar novamente. Se ainda não for possível um implante satisfatório, o dispositivo deve ser recapturado e substituído por um novo (o texto sublinhado e em itálico identifica o aviso alterado)• Os eventos adversos foram actualizados para incluir mais dados sobre o evento raro de erosão, incluindo a taxa de erosão tecidual entre 1 e 3 por cada 1000 doentes



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ENCERRAMENTO DE COMUNICAÇÃO INTERAURICULAR:

“Cirurgia deve ser considerada nos casos de maior complexidade para encerramento percutâneo”



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O QUE HÁ DE NOVO NO ENCERRAMENTO DE FORAMEN OVALE PATENTE

Factos:

- Mecanismo de AVC paradoxal através do FOP (+/- FR anatómicos) está comprovado com inúmeros casos de relação causal evidente
- **25% da população mundial** é potencial candidata a encerramento FOP (?)
- Técnica percutânea de simples execução e com taxa de sucesso alta (?)
- **1.000.000** de encerramentos de FOPs a nível mundial faz desta técnica a intervenção estrutural mais frequente em todo o mundo
- **Respect e PC Trial** não conseguiram demonstrar que o encerramento de FOP em doentes após AVC criptogénico reduz o risco de novo AVC (?)



O QUE HÁ DE NOVO NO ENCERRAMENTO DE FORAMEN OVALE PATENTE: RESPECT TRIAL



The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Closure of Patent Foramen Ovale versus Medical Therapy after Cryptogenic Stroke

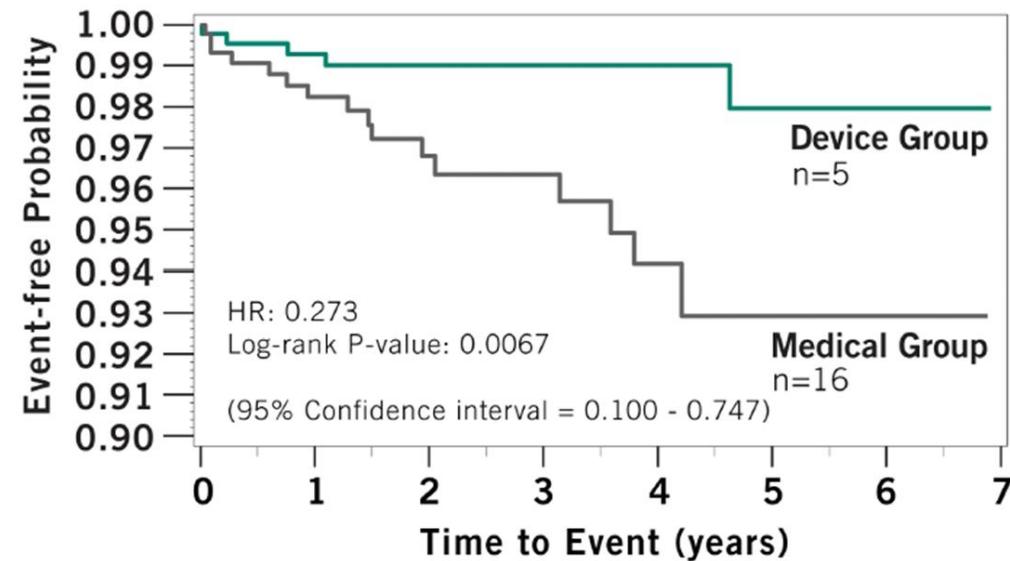
John D. Carroll, M.D., Jeffrey L. Saver, M.D., David E. Thaler, M.D., Ph.D.,
Richard W. Smalling, M.D., Ph.D., Scott Berry, Ph.D., Lee A. MacDonald, M.D.,
David S. Marks, M.D., and David L. Tirschwell, M.D.,
for the RESPECT Investigators*

n engl j med 368;12 nejm.org march 21, 2013



O QUE HÁ DE NOVO NO ENCERRAMENTO DE FORAMEN OVALE PATENTE: RESPECT TRIAL

Inc	Pri	Pr	S
-	-	-	Primary Endpoint Analysis – As Treated Cohort
-	-	-	72.7% risk reduction of stroke in favor of device
-	-	-	Device Group n=5
-	-	-	Medical Group n=16
-	-	-	(95% Confidence interval = 0.100 - 0.747)
-	-	-	HR: 0.273 Log-rank P-value: 0.0067
-	-	-	Event-free Probability
-	-	-	Time to Event (years)
-	-	-	Pa
-	-	-	Ev

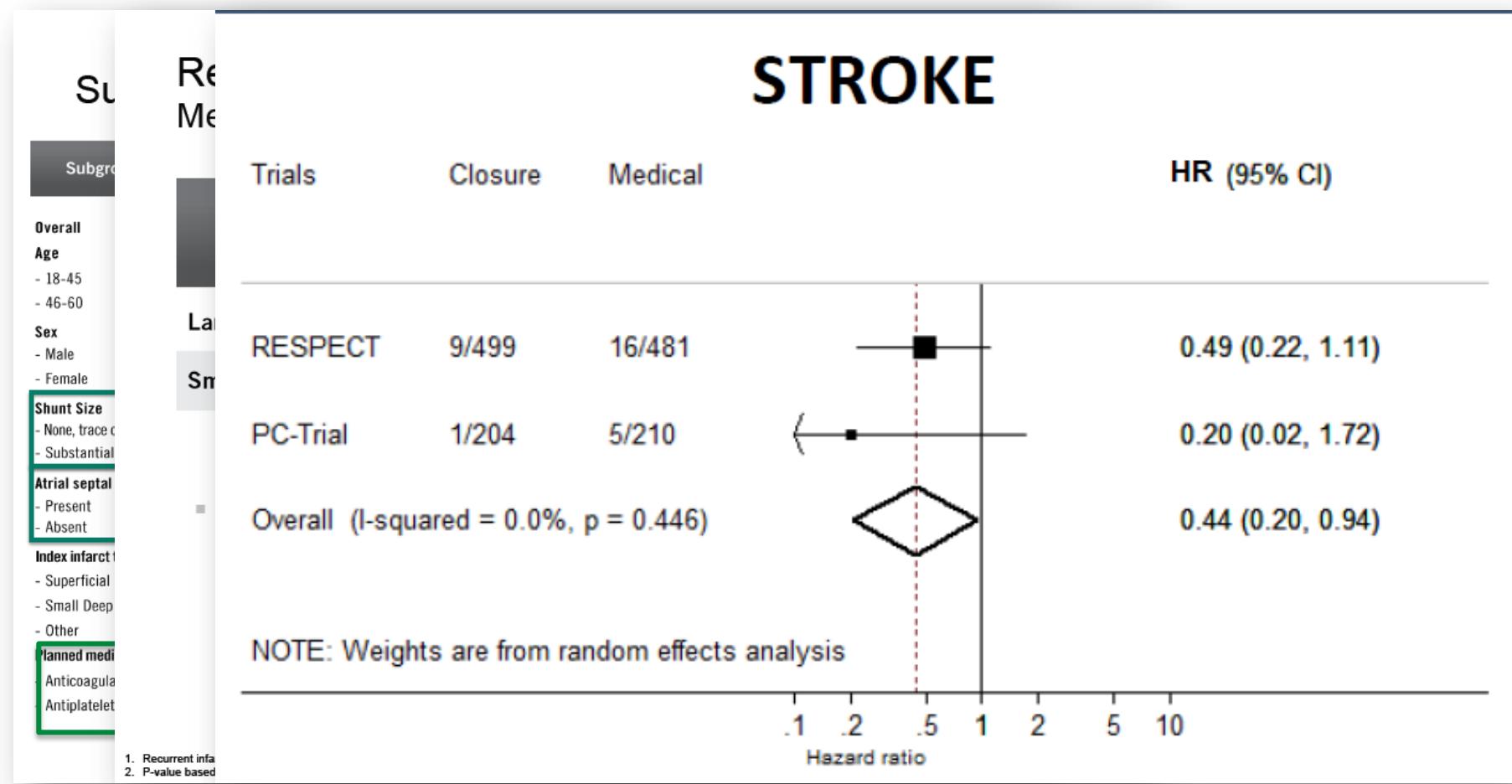


- The As Treated (AT) cohort demonstrates the treatment effect by classifying subjects into treatment groups according to the treatment actually received, regardless of the randomization assignment

22

1. Aspirin + c 1. 1. Cox model used for analysis

O QUE HÁ DE NOVO NO ENCERRAMENTO DE FORAMEN OVALE PATENTE: RESPECT TRIAL



ENCERRAMENTO PERCUTÂNEO DE FOP APÓS AVC CRIPTOGÉNICO



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Críticos em relação ás propostas de encerramento (relação causal óbvia ou muito provável)

Devemos procurar mecanismo embólico paradoxal (Factores de Risco, patologia venosa, circunstancias do AVC/AIT)

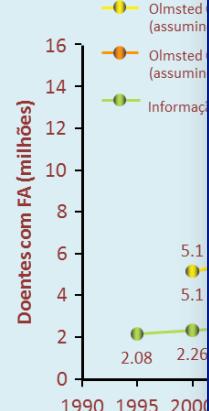


O QUE HÁ DE NOVO NO ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

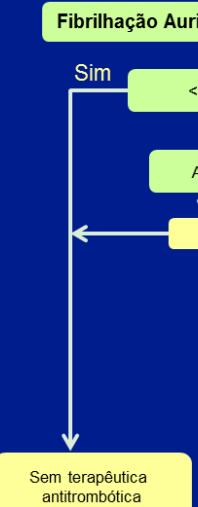
Fibrilhação auricular – “a epidemia do século XXI”

PREVA

Aumento



Guidelines I



I. AL, et al. J Am Coll Cardiol 2005

Camm AJ et al. Eur Heart J 2012

- Na prática clínica, os doentes com FA de elevado risco...

Doentes com FA de elevado risco

50% tratados com ACO

50% param ACO entre 3-5 anos

50% mantêm INR adequados

50% não tratados com ACO

1. Miyasaka Y et al. Circulation 2006;114:119–125;

*. Go AS et al. JAMA 2001;285:2370–2375; 2. Heeringa J et al. Eur Heart J 2006;27:949–953; 3. Frost L et al. Int J Cardiol 2005;103:78–84; 4. DeWilde S et al. Heart 2006;92:1064–1070; 5. Miyasaka Y et al. Circulation 2006;114:119–125; 6. Zhou Z and Hu D. J Epidemiol 2008;18:209–216; 7. Fuster V et al. Circulation 2006;114:700–752; 8. Zimmerman LI et al. Arq Bras Cardiol 2009;92:1–39; 9. ESC Guidelines Eur Heart J 2010; 31:2369–2429; 10. Naccarelli GV et al. Am J Cardiol 2009;104(11):1534–9; 11. Daniel et al. Estudo FAMA, Rev Port Cardiol, vol 29 Mar 2010.



*Paradoxo do idoso:
«maior eficácia mas maior risco»*



ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Novos anticoagulantes no tratamento da FA



Systematic Review and Adjusted Indirect Comparison Meta-Analysis of Oral Anticoagulants in Atrial Fibrillation

William L. Baker and Olivia J. Phung

Circ Cardiovasc Qual Outcomes. 2012;5:711-719; originally published online August 21, 2012;
doi: 10.1161/CIRCOUTCOMES.112.966572

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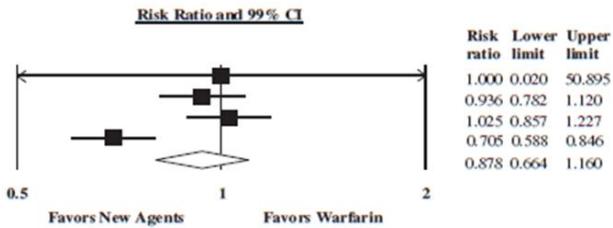


ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Novos anticoagulantes no tratamento da FA

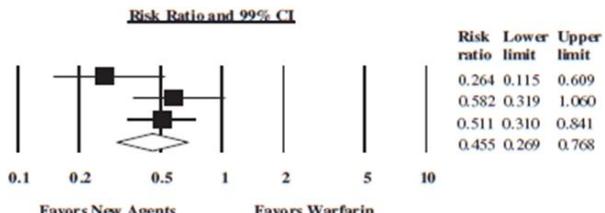
F Major Bleed

Study Name	Events / Total	
	New Agents	Warfarin
PETRO, 2007	0 / 100	0 / 70
RE-LY, 2009	375 / 6076	397 / 6022
ROCKET-AF, 2011	395 / 7111	386 / 7125
ARISTOTLE, 2011	327 / 9088	462 / 9052
TOTAL	1097 / 22375	1245 / 22269
p = 0.23		
F = 80,6%		
Egger p = 0.98		



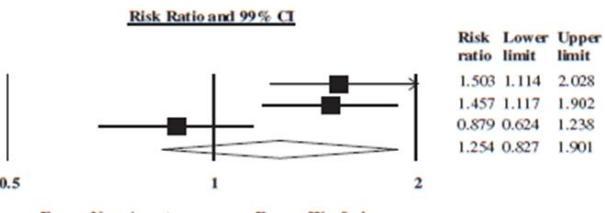
G Hemorrhagic Stroke

Study Name	Events / Total	
	New Agents	Warfarin
RE-LY, 2009	12 / 6076	45 / 6022
ROCKET-AF, 2011	29 / 7061	50 / 7082
ARISTOTLE, 2011	40 / 9120	78 / 9081
TOTAL	81 / 22257	179 / 22185
p = 0.0001		
F = 52,1%		
Egger p = NA		



H Gastrointestinal Bleed

Study Name	Events / Total	
	New Agents	Warfarin
RE-LY, 2009	182 / 6076	120 / 6022
ROCKET-AF, 2011	224 / 7111	154 / 7125
ARISTOTLE, 2011	105 / 9088	119 / 9052
TOTAL	511 / 22275	393 / 22199
p = 0.16		
I² = 82,5%		
Egger p = NA		



A incidencia de Hemorragias major no conjunto de 4 estudos Randomizados (44.644 dts) mostrou:

- Incidencia de hemorragias major oscila entre **3,5%** (grupo Apixaban do estudo ARISTOTELES e **6,5%** do grupo Varfine do estudo RELY)
- No conjunto de dts nestes studos verificaram-se 2342 Hemorragias major ou seja **5,2%**

ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Que fazer aos doentes que fazem ACO e tem hemorragia?



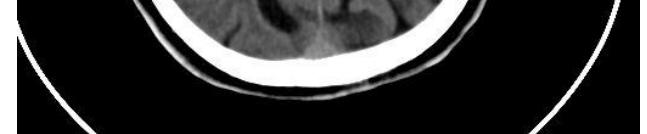
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Onde estão estes doentes ?
Só em Portugal deverão ser cerca de
5000 dts



O que podemos fazer por
estes doentes ?





Management of Atrial Fibrillation Focus of 2012 Update

- Anticoagulation risk stratification
- Use of novel oral anticoagulants (NOACs)
- Left atrial appendage occlusion/excision
- Pharmacological cardioversion (vernakalant)

Recomendação para os doentes com contra-indicação aos ACO ou elevado risco hemorrágico

www.escardio.org/guidelines

European Heart Journal 2012;33:2719-2747 -
doi:10.1093/eurheartj/ehs253



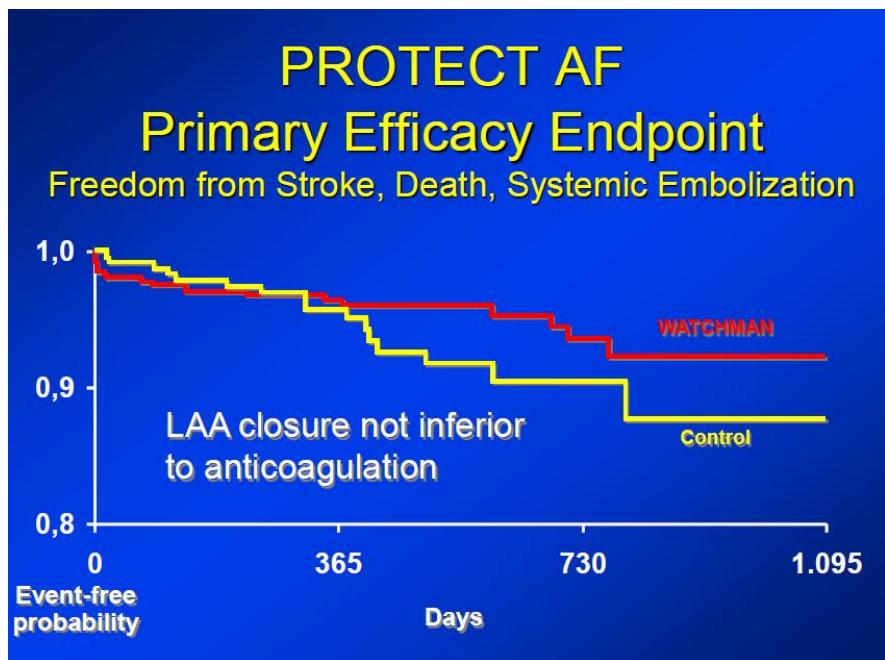
ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Evidência clínica



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PROTECT AF



PROTECT AF: The Mortality Effects of LAA Closure vs. Warfarin for Stroke Prophylaxis in A-fib

Four-year results for patients with nonvalvular A-fib randomized to percutaneous LAA closure with Watchman ($n = 463$) or warfarin alone ($n = 244$).

- At 4 years, the primary efficacy event rate per 100 patient years was lower with Watchman (2.3%) than controls (3.8%; RR 0.60; 95% CI 0.41-1.05)
- Compared with warfarin, Watchman was linked with lower risk of all-cause (3.2% vs. 4.8%; HR 0.66; $P = 0.0379$) and cardiovascular mortality (1.0% vs. 2.4%; HR 0.40; $P = 0.0045$)
- Efficacy results confirmed through intention-to-treat, post-procedure, per-protocol, and terminal therapy analyses

Conclusion: For patients with nonvalvular A-fib, closing the LAA with the Watchman device has been shown to be superior to warfarin alone.

Reddy VY. Heart Rhythm 2013;
Denver, CO.



The Source for Interventional Cardiovascular News and Education



ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Evidência clínica



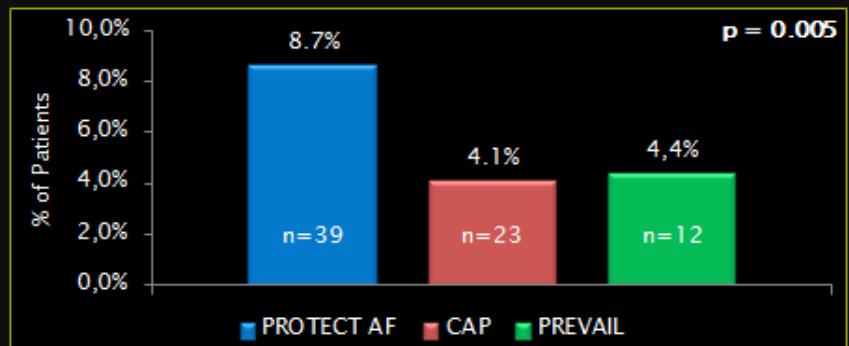
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PROTECT AF /CAP /PREVAIL (Safety Endpoint)

Vascular Complications

7 Day Serious Procedure/Device Related

- Composite of vascular complications includes cardiac perforation, pericardial effusion with tamponade, ischemic stroke, device embolization, and other vascular complications¹

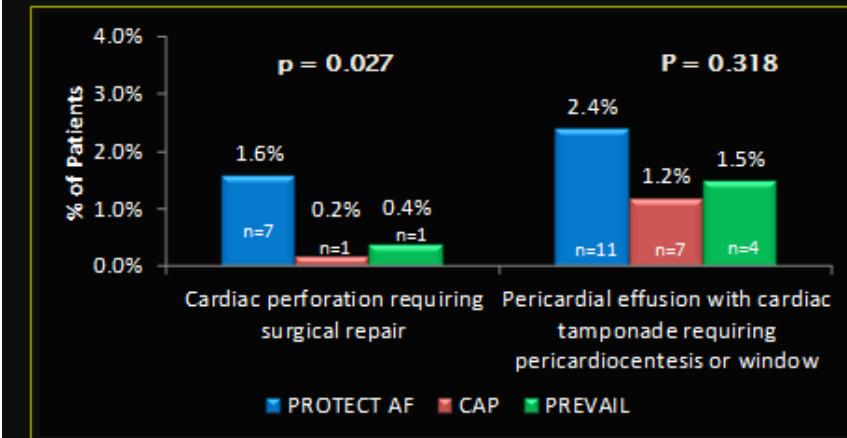


No procedure-related deaths reported in any of the trials

PROTECT-AF and CAP data from Reddy VY et al. Circulation. 2011;123:417-424.
Includes observed PE not necessitating intervention, AV fistula, major bleeding requiring transfusion, pseudotamponade, hematoma and groin bleeding.

Caution: In the United States, WATCHMAN is an investigational device limited by Federal law and investigational use only. Not for sale in the US. Prior to use please review device indications, contraindications, warnings, precautions, adverse events, and operational instructions. Only available according to applicable local law. CE Mark received in 2005

Pericardial Effusions Requiring Intervention



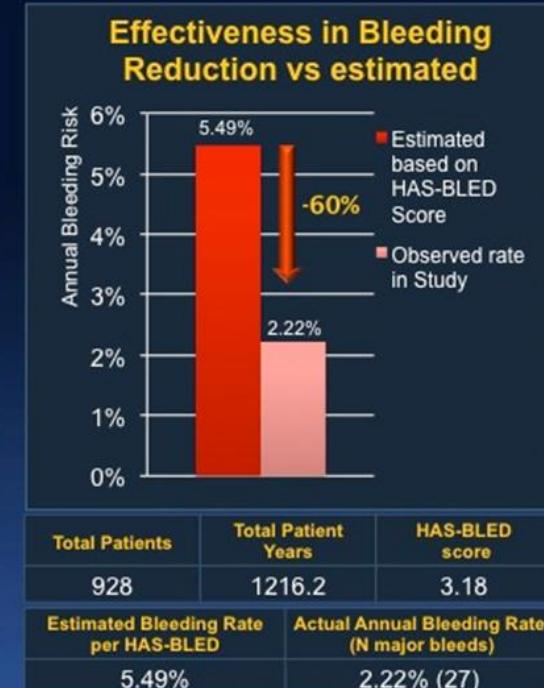
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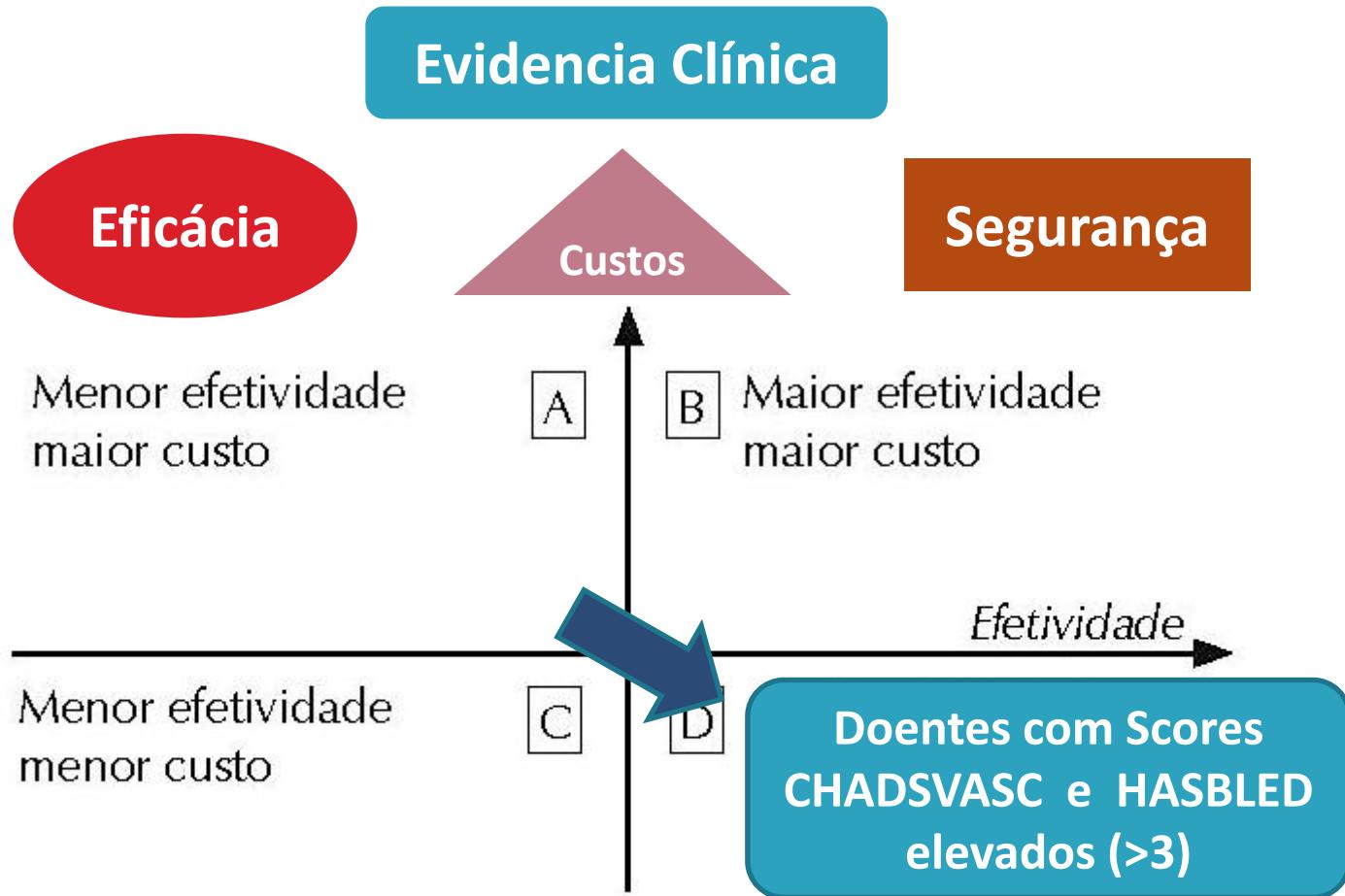
O QUE HÁ DE NOVO NO ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

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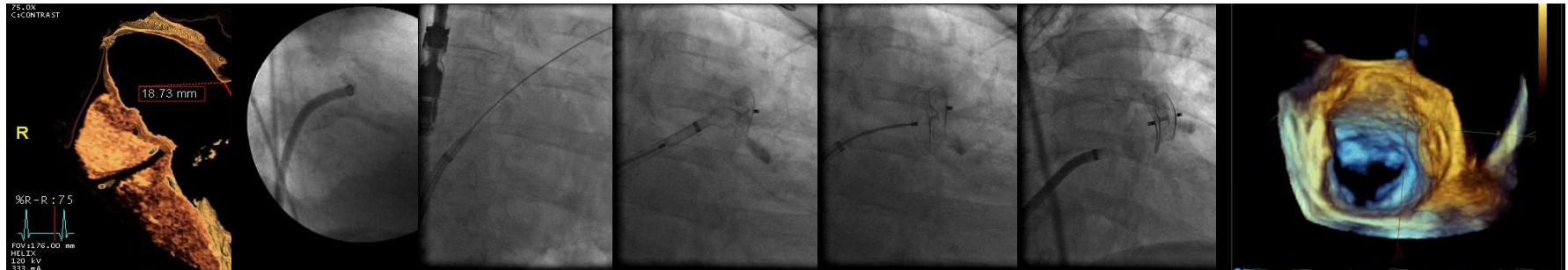
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ENCERRAMENTO DE APÊNDICE AURICULAR ESQUERDO

Doentes para Referenciar

Doentes com FA não valvular com ChadsVasc2 de 1+

- Evento hemorrágico relacionado com Varfine (INR <3) ou com NOACs
- Risco hemorrágico elevado que não consideramos á partida para ACO (HasBled de 3+, co-medicação, anemia crónica, má compliance ...)
- AVC com trombo identificado no AAE a tomar ACO (com INR >2)
(...)



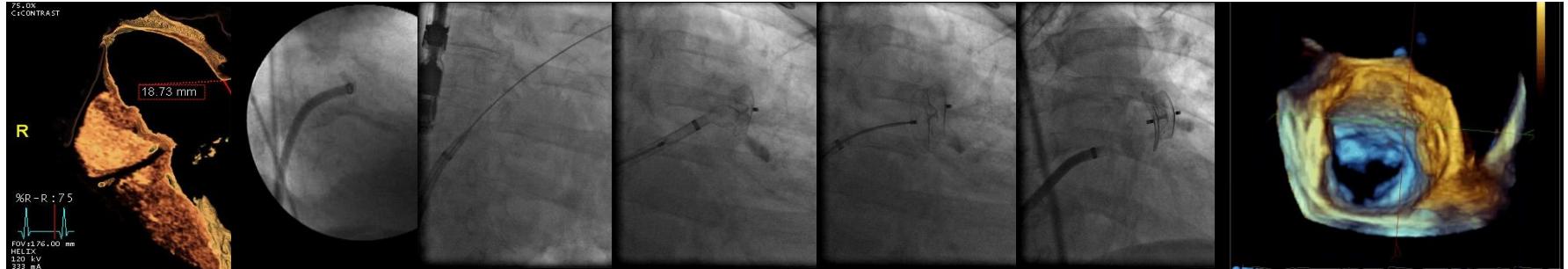


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(...)



Em medicina, como em tudo o resto, aplica-se a máxima
“o que é verdade hoje pode não ser amanhã”

