

PHAR 451

Ischemic Stroke Prevention in patients with Atrial Fibrillation

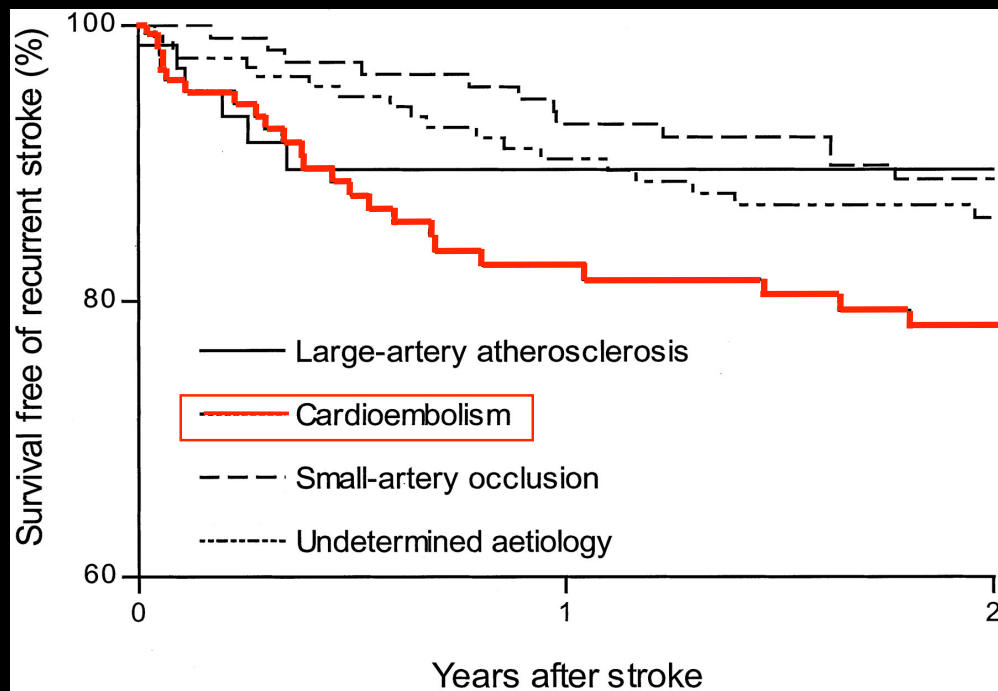


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University of British Columbia

Objective

After the session, and upon personal reflection & study, students will be able to DESIGN and RATIONALIZE using EVIDENCE, a stroke prevention regimen (drugs, doses, routes, frequency) for a patient with atrial fibrillation which incorporates their stroke risk, bleeding risk, access to coverage, personal values, and ability to be monitored.

2-Year Stroke-free Survival After Ischemic Stroke



Kolominsky-Rabas et al. Stroke 2001;32:2735-40

CASE

- 78 y/o F with newly-identified asymptomatic atrial fibrillation
- PMH:
 - DM2 (diet-controlled)
 - TIA in 1999 (receiving ASA 80mg/d since)
 - MI in 2009



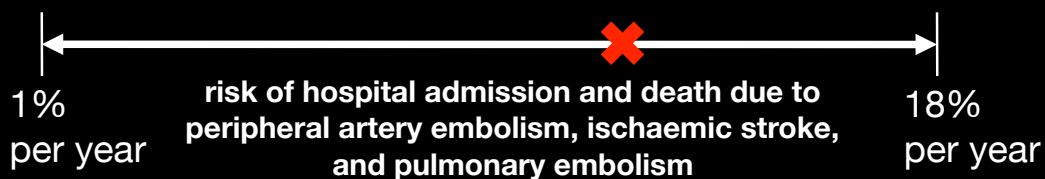
What is the most appropriate antithrombotic therapy for stroke prophylaxis in this patient?

Stroke risk in chronic Atrial Fib

"CHADS2"

LV Dysfunction (CHF)
 HTN
 Age > 75 1
 Diabetes 1
 Previous Stroke/TIA 2
 CHADS2 score: 4

CHADS2 Score	Stroke risk/ year
0	1.2%
1	3.6%
2	5.4%
3	9.9%
4	13.7%
5	12.6%
6	17.2%



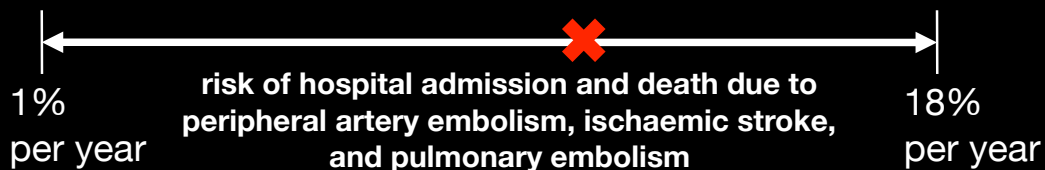
Olesen JB, et al. BMJ 2011;342:d124

Stroke risk in chronic Atrial Fib

"CHA2DS2-VASc"

LV Dysfunction (CHF)
 HTN
 Age > 75 2
 Diabetes 1
 Previous Stroke/TIA 2
 Prior MI, peripheral artery disease, or aortic plaque 1
 Age 65-75
 Female 1
 CHA2DS2-VASc score: 7

CHADS2 Score	Stroke risk/ year
0	0.7%
1	1.5%
2	2.9%
3	4.3%
4	6.5%
5	10%
6	12.5%
7	14%
8	14.1%
9	16%



Olesen JB, et al. BMJ 2011;342:d124

SPARC - Stroke Prevention in Atrial Fibrillation Risk Tool

for estimating risk of stroke and benefits & risks of antithrombotic therapy in patients with chronic atrial fibrillation

[references/notes](#)

version 7, January 2015

Developed by Peter Loewen, ACPR, Pharm.D., FCSHP

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In your patient with atrial fibrillation, which of the following stroke or bleeding risk factors are present?

CHADS2 CRITERIA

- CHF/LV dysfunction (diagnosed at any time in the past)
- Hypertension (controlled or uncontrolled)
- Age > 75
- Diabetes Type I or II (controlled or uncontrolled)
- TIA or stroke (at any time in the past)

CHADS2 SCORE (0-6):0

CHA2DS2-VASc CRITERIA

- Prior MI, peripheral artery disease, or aortic plaque
- Age 65-75
- Female

CHA2DS2-VASc SCORE (0-9):0

HAS-BLED CRITERIA*

- Abnormal renal function (dialysis, SCr>200 mmol/L, or transplant)
- Abnormal liver function (cirrhosis or liver enzymes >3x ULN)
- History of major bleeding (any cause)
- History of labile INR (time in therapeutic range <60%)
- Current "excess" use of alcohol
- Currently taking antiplatelet drug(s) or NSAID(s)

HAS-BLED SCORE (0-9)*:0

www.sparctool.com

www.sparctool.com

<https://itunes.apple.com/ca/app/anticoagevaluator/id609795286?mt=8>

Stroke Prevention in AF

What we do	In whom	Effect vs. placebo	Source
ASA	CHADS2 0-1 <u>OR</u> unable to anticoagulate	RR 0.78	Ann Intern Med. 2007;146:857-867
warfarin	CHADS2 > 1	RR 0.33	
dabigatran	CHADS2 > 1 & if preferred over warfarin, labile INR, coverage available	110mg bid: similar efficacy+less bleeding vs. warfarin	RE-LY
rivaroxaban		150mg bid: superior efficacy +similar bleeding vs. warfarin	ROCKET-AF
apixaban		superior efficacy and safety to warfarin	ARISTOTLE
(edoxaban)	not marketed in Canada yet	efficacy similar to warfarin, less bleeding	ENGAGE-AF-TIMI48

EASY THING TO REMEMBER:

WARFARIN in Atrial Fibrillation

RELATIVE RISK OF
STROKE = 0.33
vs. no therapy

Hart et al. Ann Intern Med 1999;131:492-501
Ann Intern Med. 2007;146:857-867

EASY THING TO REMEMBER:

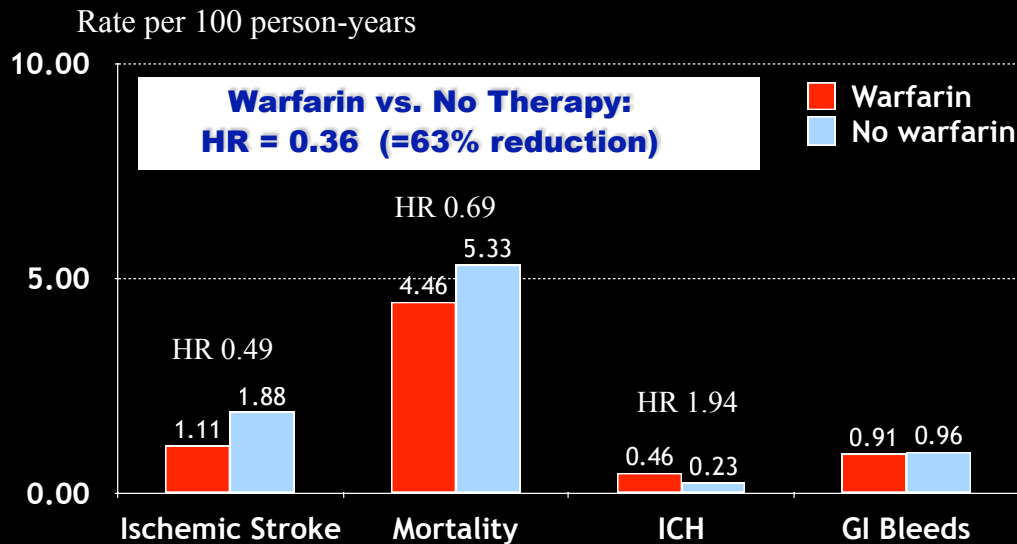
ASPIRIN in Atrial Fibrillation

RELATIVE RISK OF
STROKE = 0.78
vs. no therapy

Hart et al. Ann Intern Med 1999;131:492-501
Ann Intern Med. 2007;146:857-867

Effectiveness of Warfarin in AF

- Cohort study, N=11,526 wth AF, mean 71 y/o
- 2.2 years of observation



Go et al. JAMA 2003;290:2685-92

Early release, published at www.cmaj.ca on November 26, 2012. Subject to revision.

CMAJ

RESEARCH

Rates of hemorrhage during warfarin therapy for atrial fibrillation

Tara Gomes MHS, Muhammad M. Mamdani PharmD MPH, Anne M. Holbrook MD PharmD, J. Michael Paterson MSc, Chelsea Hellings MSc, David N. Juurlink MD PhD

ABSTRACT

Background: Although warfarin has been extensively studied in clinical trials, little is known about rates of hemorrhage attributable to its use in routine clinical practice. Our objective was to examine incident hemorrhagic events in a large population-based cohort of patients with atrial fibrillation who were starting treatment with warfarin.

Methods: We conducted a population-based cohort study involving residents of Ontario (age ≥ 66 yr) with atrial fibrillation who started taking warfarin between Apr. 1, 1997, and Mar. 31, 2008. We defined a major hemorrhage as any visit to hospital for hemorrhage. We determined crude rates of hemorrhage during warfarin treatment, overall and stratified by CHADS₂ score (congestive heart failure, hypertension, age ≥ 75 yr, diabetes mellitus and prior stroke,

rate of hemorrhage was 3.8% (95% confidence interval [CI] 3.8%–3.9%) per person-year. The risk of major hemorrhage was highest during the first 30 days of treatment. During this period, rates of hemorrhage were 11.8% (95% CI 11.1%–12.5%) per person-year in all patients and 16.7% (95% CI 14.3%–19.4%) per person-year among patients with a CHADS₂ score of 4 or greater. Over the 5-year follow-up, 10 840 patients (8.7%) visited the hospital for hemorrhage; of these patients, 1963 (18.1%) died in hospital or within 7 days of being discharged.

Interpretation: In this large cohort of older patients with atrial fibrillation, we found that rates of hemorrhage are highest within the first 30 days of warfarin therapy. These rates are considerably higher than the rates of 1%–3% reported in randomized controlled trials

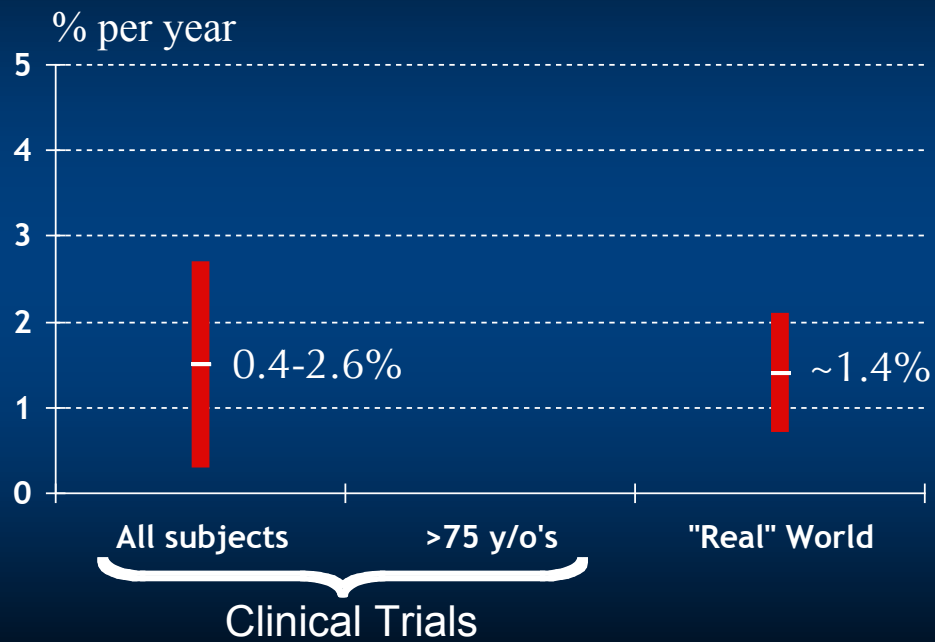
Competing interests: Tara Gomes, Chelsea Hellings and David Juurlink have received grant funding from the Ontario Drug Policy Research Network. Muhammad Mamdani is a consultant for AstraZeneca, Bristol-Myers Squibb, Eli Lilly, GlaxoSmithKline, Hoffmann-La Roche, Novartis, Novo Nordisk and Pfizer. No other competing interests were declared.

This article has been peer reviewed.

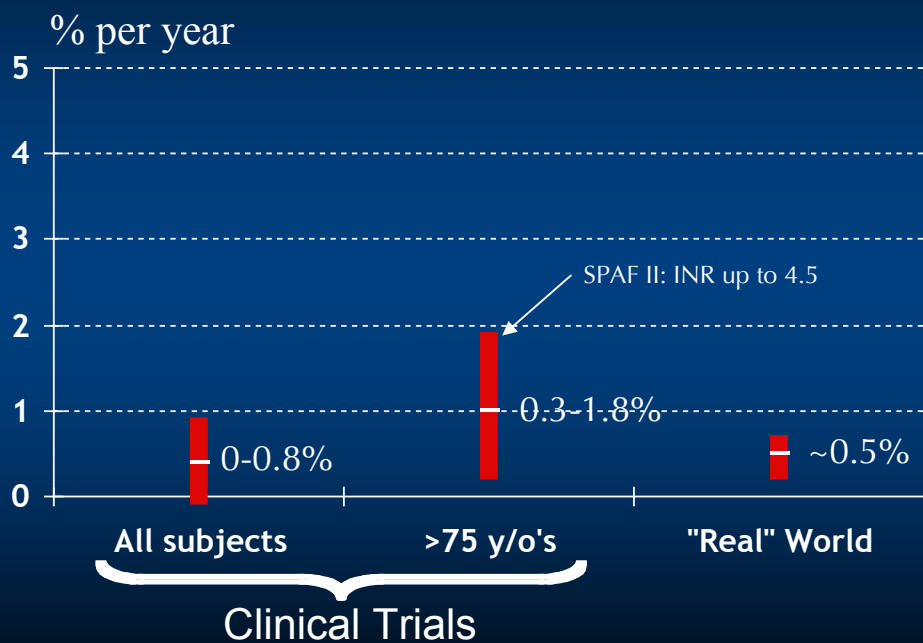
Correspondence to: Tara Gomes, tara.gomes@ices.on.ca

CMAJ 2012. DOI:10.1503/cmaj.121218

Major Bleeding with warfarin

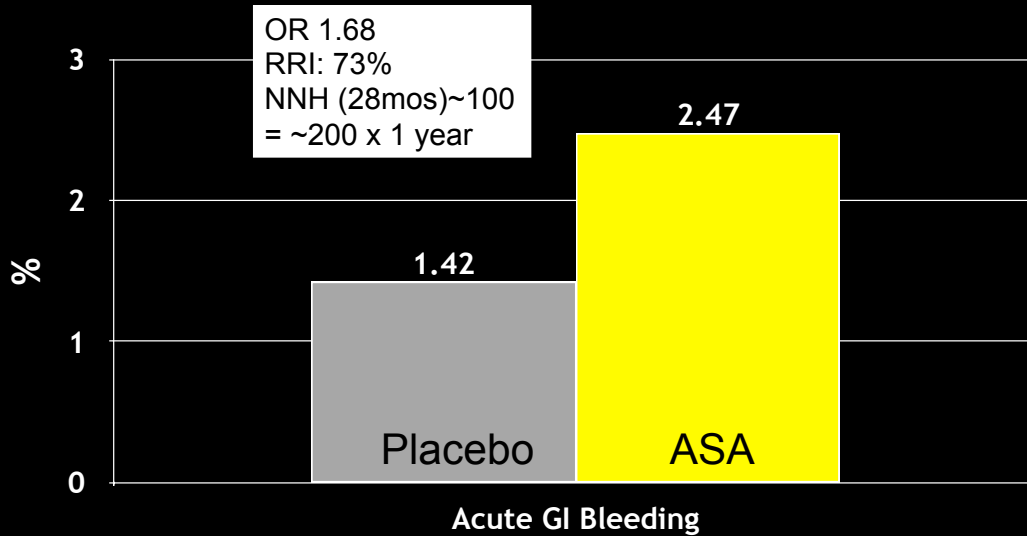


ICH with warfarin



Safety of ASA: Acute GI Bleeds

N=24 trials, 66,000 patients. Average follow-up 28 mos.



Derry & Loke. BMJ 2000;321:1183-7
consistent with ATTC 2009; Lancet 2009; 373: 1849-60
Lanas et al. Clin Gastro Hepatol 2011;9:762

Safety of low-dose ASA: Bleeding

N=35 trials, 87,581 patients. 338,735 person-years followup. Average follow-up 3.9 y.

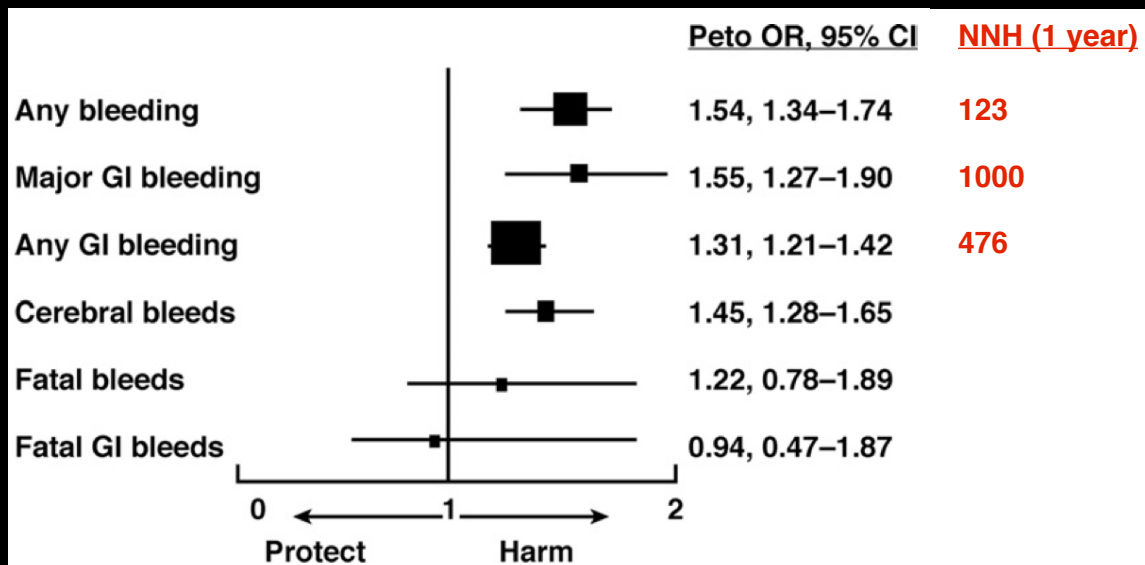
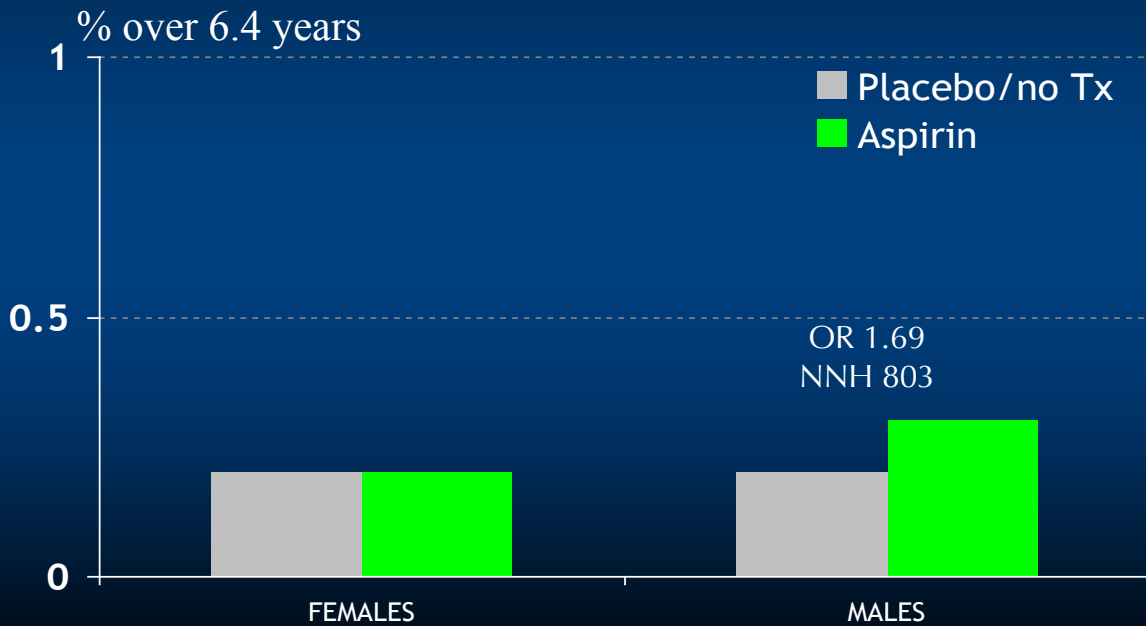


Figure 1. Bleeding events associated with low-dose ASA alone vs controls.

Safety of ASA: Intracranial Hemorrhage

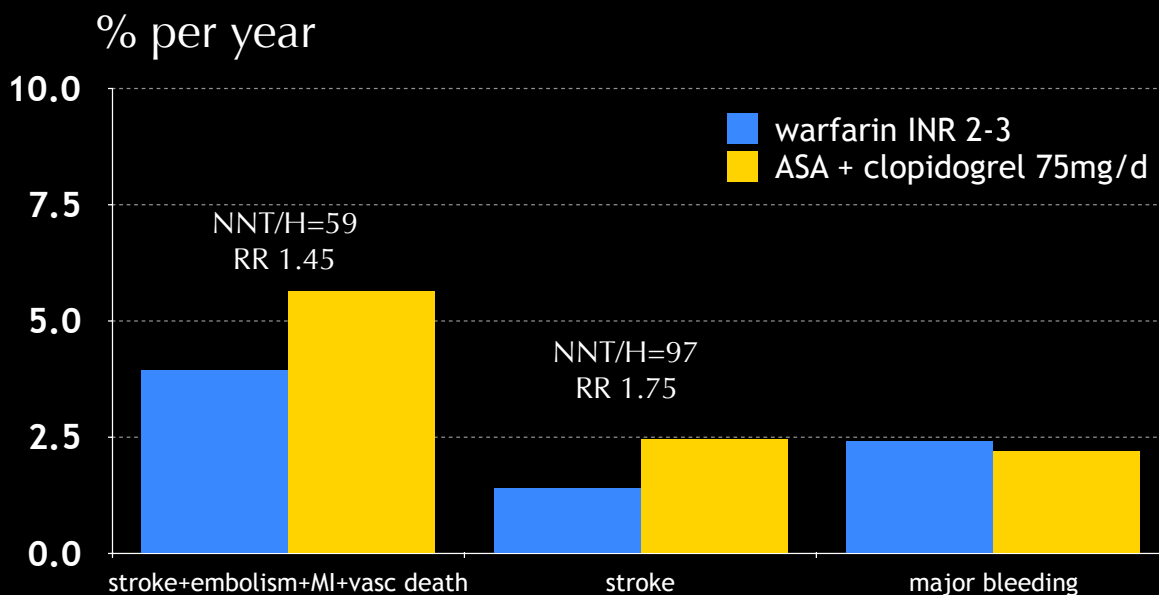
N=6 trials. 51,342 women, 44,114 men.
ASA 75mg - 500 mg/d, mean 6.4 years followup.



Berger JS et al. JAMA 2006;295:306-13

ACTIVE W

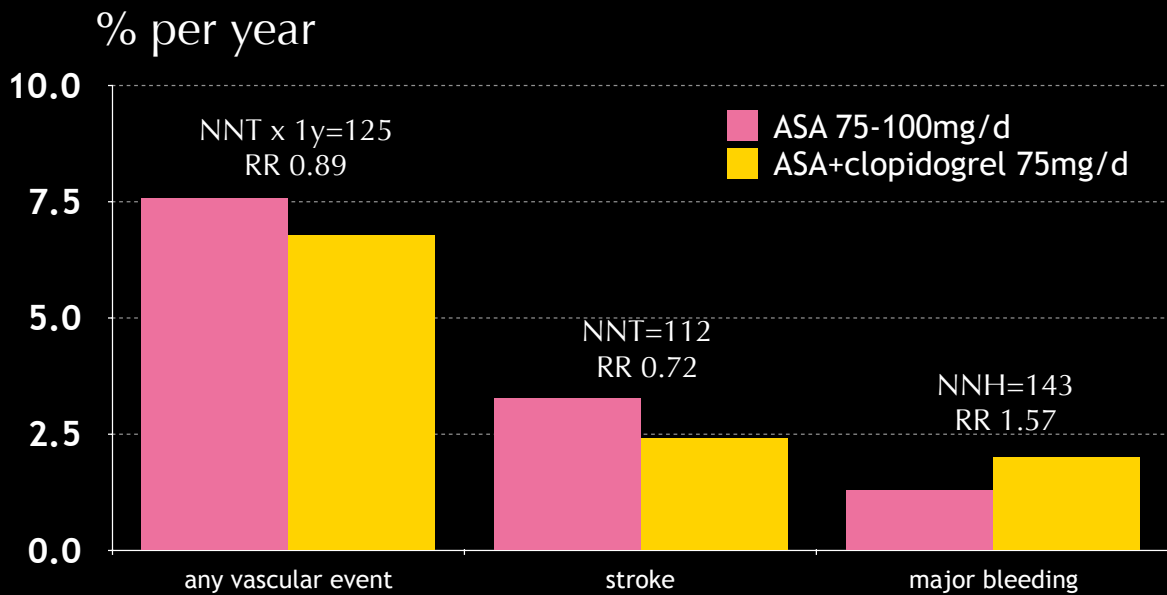
N= 6,707 AF patients with 2+ risk factors for stroke (CHADS + PVD), felt to be candidates for warfarin therapy. 33% >75 y/o. Stopped early.



ACTIVE-W. Lancet 2006;367:1903-12

ACTIVE A

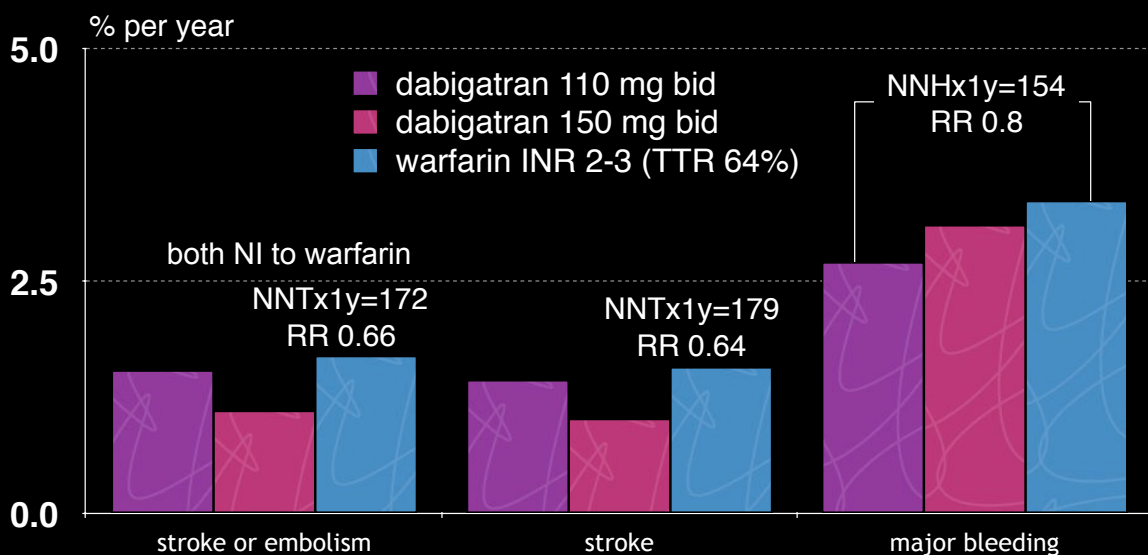
N= 7,554 AF patients with 1+ risk factors for stroke (CHADS₂+CAD), felt to be "unsuitable" for warfarin therapy. Median 3.6y followup.



ACTIVE-A. NEJM 2009;360 [1APR09]

RE-LY: dabigatran vs. warfarin in AF

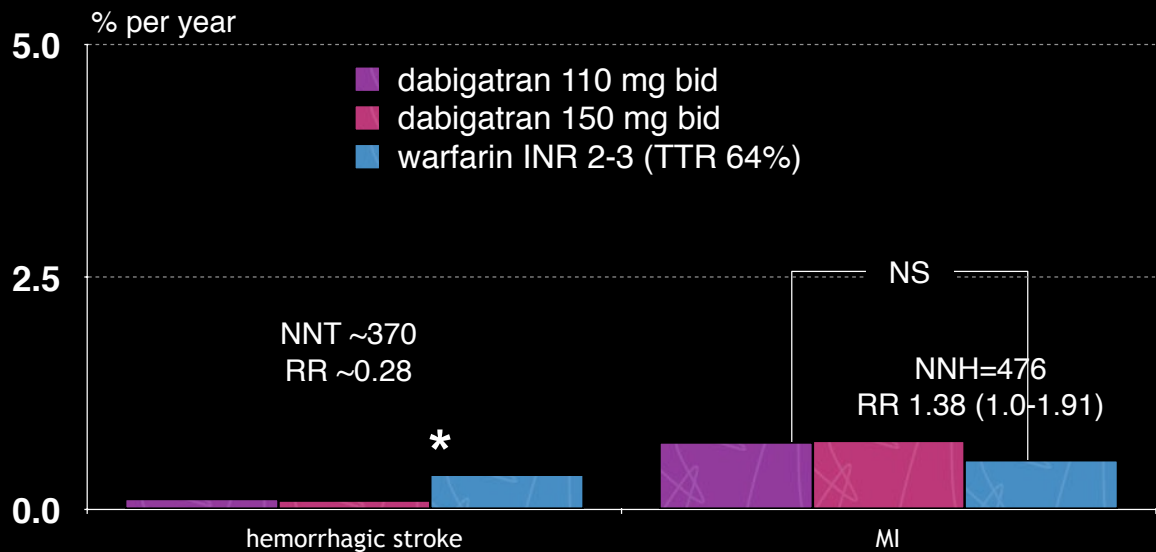
N= 18,113 AF patients with 1+ risk factors for stroke (mean CHADS₂ score 2.1). Median 2y followup. Non-inferiority trial.



RE-LY. NEJM 2009;361

RE-LY: dabigatran vs. warfarin in AF - other endpoints

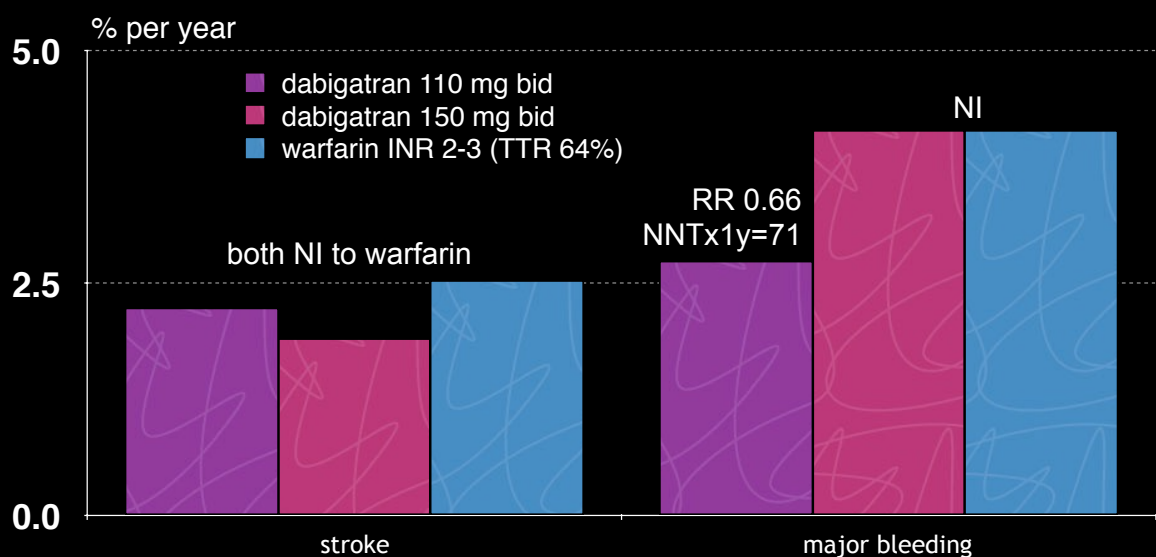
N= 18,113 AF patients with 1+ risk factors for stroke (mean CHADS2 score 2.1). Median 2y followup. Non-inferiority trial.



RE-LY. NEJM 2009;361

RE-LY: dabigatran vs. warfarin in AF - SECONDARY prevention

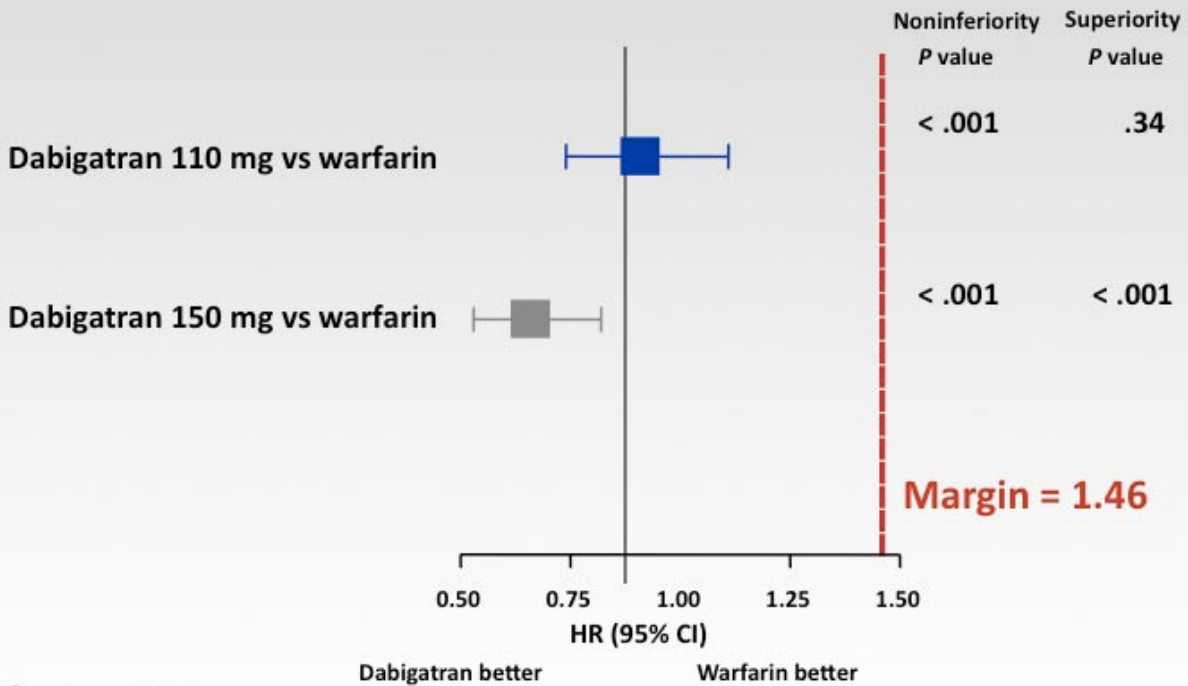
N= 3,623 AF patients with prior stroke/TIA. Median 2y followup. Pre-specified secondary analysis



RE-LY. Lancet Neurol 2010; 9: 1157-63

RE-LY[®] Trial

Primary Endpoints: All Strokes and Systemic Embolic Events



Connolly SJ, et al. *N Engl J Med.* 2009;361:1139-1151.



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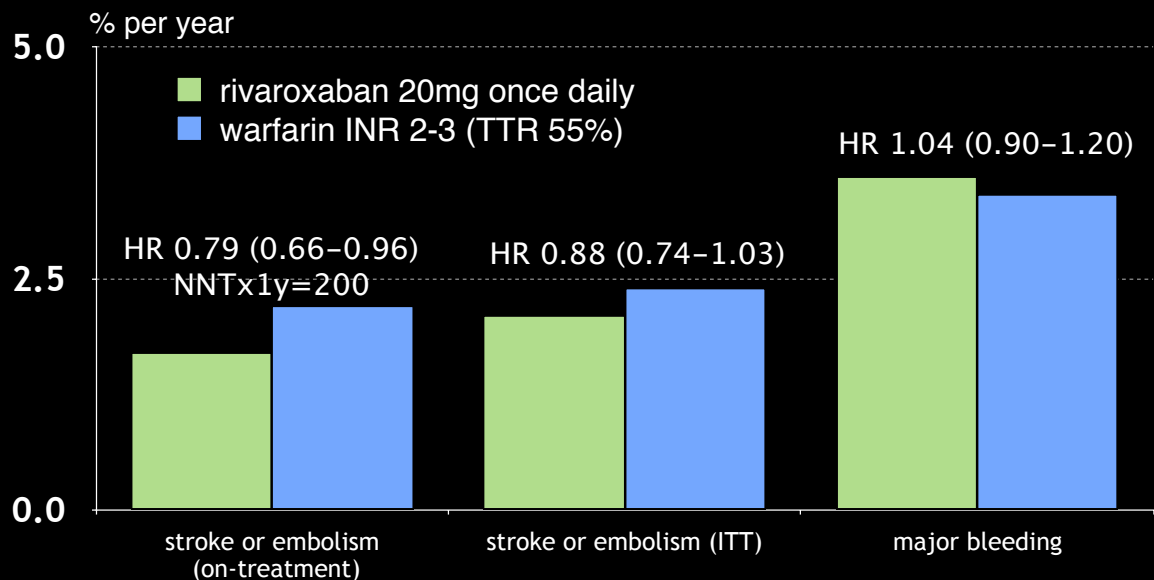
- Safety Information
- Safety Alerts for Human Medical Products
- 2012 Safety Alerts for Human Medical Products
- 2011 Safety Alerts for Human Medical Products

Pradaxa (dabigatran etexilate mesylate): Drug Safety Communication - Safety Review of Post-Market Reports of Serious Bleeding Events

UPDATED 11/02/2012. The FDA evaluated new information about the risk of serious bleeding associated with use of the anticoagulants (blood thinners) dabigatran (Pradaxa) and warfarin (Coumadin, Jantoven, and generics). This assessment was done using insurance claims and administrative data from FDA's Mini-Sentinel pilot of the Sentinel Initiative. The results of this assessment indicate that bleeding rates associated with new use of Pradaxa do not appear to be higher than bleeding rates associated with new use of warfarin, which is consistent with observations from the large clinical trial used to approve Pradaxa (the RE-LY trial). FDA is continuing to evaluate multiple sources of data in the ongoing safety review of this issue. See the Data Summary in the 11/02/2012 Drug Safety Communication below for additional information.

Rivaroxaban: ROCKET-AF

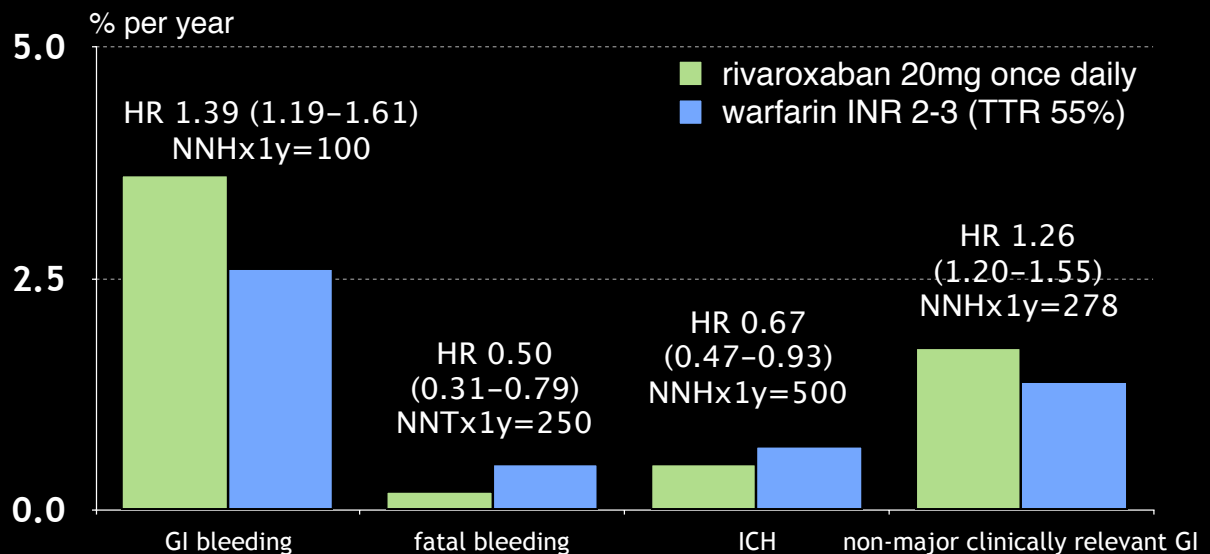
N= 14,264 AF patients, 87% with CHADS2 >2.
Median 20 months followup. DB, Non-inferiority trial.



ROCKET-AF. NEJM 2011;10.1056/NEJMoa1009638 (10AUG11)

Rivaroxaban: ROCKET-AF - bleeding breakdown

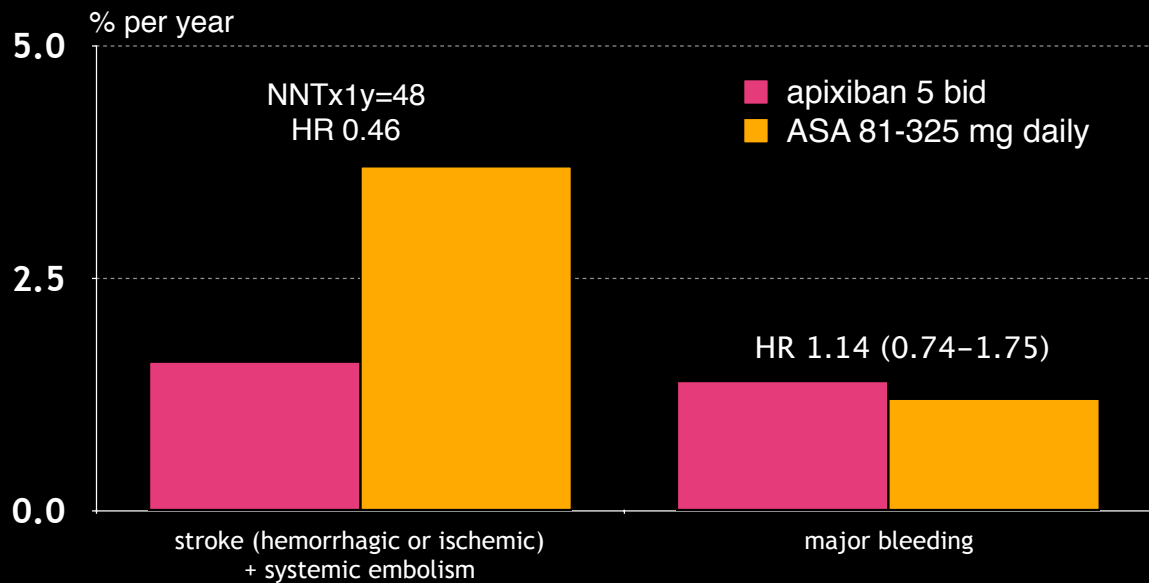
N= 14,264 AF patients, 87% with CHADS2 >2.
Median 20 months followup. DB, Non-inferiority trial.



ROCKET-AF. CHEST.October 2012;142(4_MeetingAbstracts):84A-84A

Apixaban: AVERROES

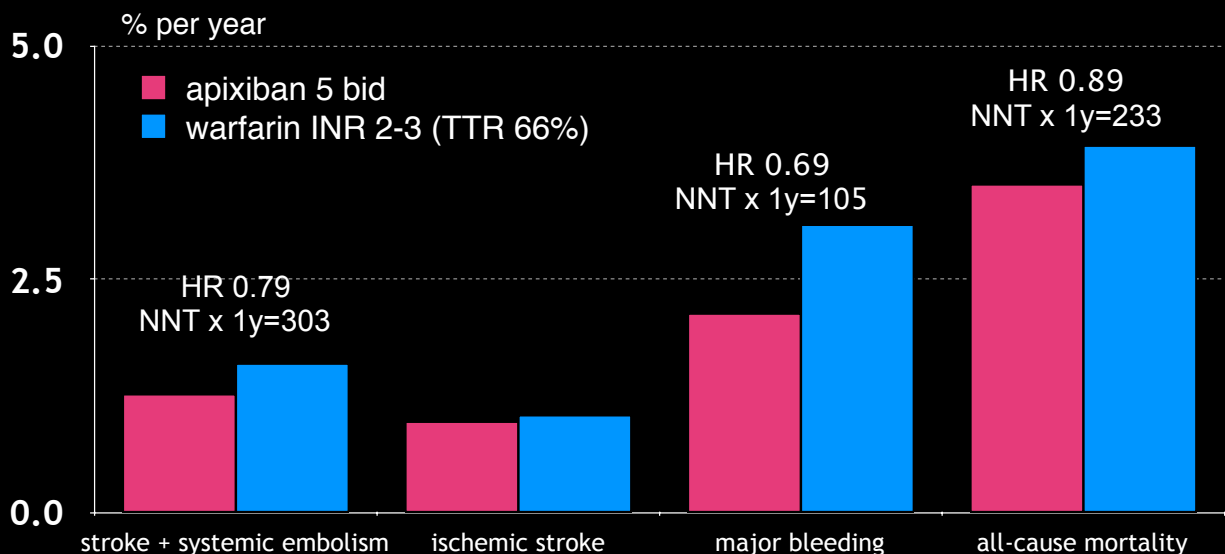
N= 5,599 AF patients with AF + intolerant or “unsuitable” for warfarin. Max 3y followup, mean 1.1 years (stopped early). Superiority trial.



AVERROES. N Engl J Med 2011 (10.1056/NEJMoa1007432)

Apixaban: ARISTOTLE

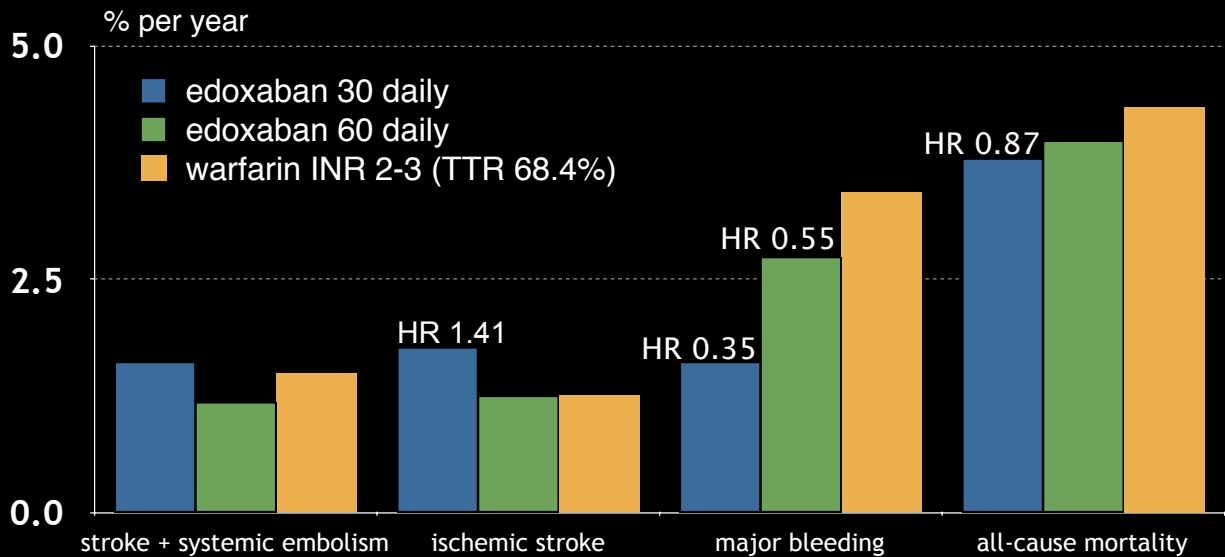
N= 18,201 AF patients with AF. ~70 had CHADS2 score >1. Median 1.8y followup. DB non-inferiority trial.



ARISTOTLE. N Engl J Med 2011. (10.1056/NEJMoa1107039) 28AUG11

Edoxaban: ENGAGE AF-TIMI 48

N= 21,105 AF patients with AF & CHADS2 score >1. Median 2.8y followup. DB non-inferiority trial.



ENGAGE AF-TIMI 48. NEJM 2013; 19NOV13

NOAC PharmaCare Coverage in BC

Dabigatran

- <http://www2.gov.bc.ca/gov/content/health/practitioner-professional-resources/pharmacare/prescribers/limited-coverage-drug-program/limited-coverage-drugs-dabigatran>

Rivaroxaban

- <http://www2.gov.bc.ca/gov/content/health/practitioner-professional-resources/pharmacare/prescribers/limited-coverage-drug-program/limited-coverage-drugs-rivaroxaban-for-atrial-fibrillation-af>

Apixaban

- <http://www2.gov.bc.ca/gov/content/health/practitioner-professional-resources/pharmacare/prescribers/limited-coverage-drug-program/limited-coverage-drugs-apixaban-for-atrial-fibrillation-af>

<http://www2.gov.bc.ca/gov/content/health/practitioner-professional-resources/pharmacare>

The “CCS Algorithm” for OAC Therapy in AF

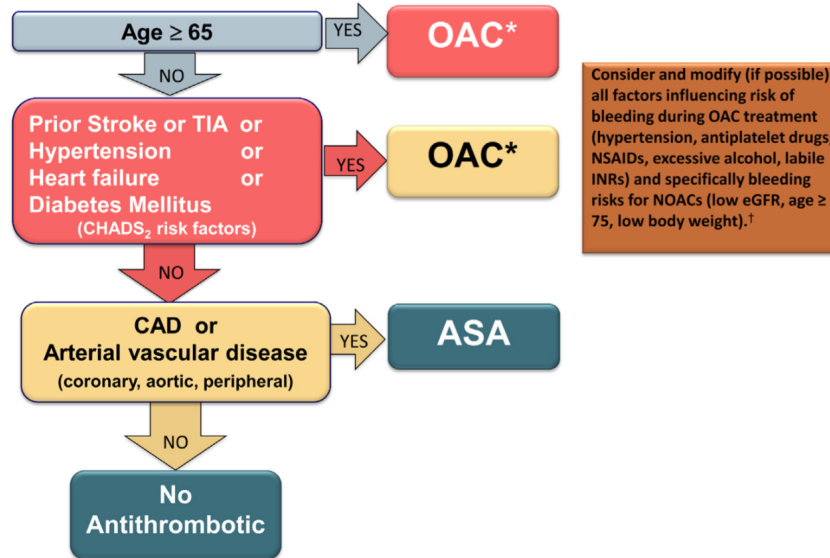


Figure 1. The simplified “CCS algorithm” for deciding which patients with atrial fibrillation (AF) or atrial flutter (AFL) should receive oral anti-coagulation (OAC) therapy. * We suggest that a NOAC be used in preference to warfarin for non-valvular AF. † Might require lower dosing. ASA, acetylsalicylic acid; CAD, coronary artery disease; CCS, Canadian Cardiovascular Society; CHADS₂, Congestive Heart Failure, Hypertension, Age, Diabetes, Stroke/Transient Ischemic Attack; eGFR, estimated glomerular filtration rate; INR, international normalized ratio; NOAC, novel oral anticoagulant; NSAID, nonsteroidal anti-inflammatory drug; TIA, transient ischemic attack.