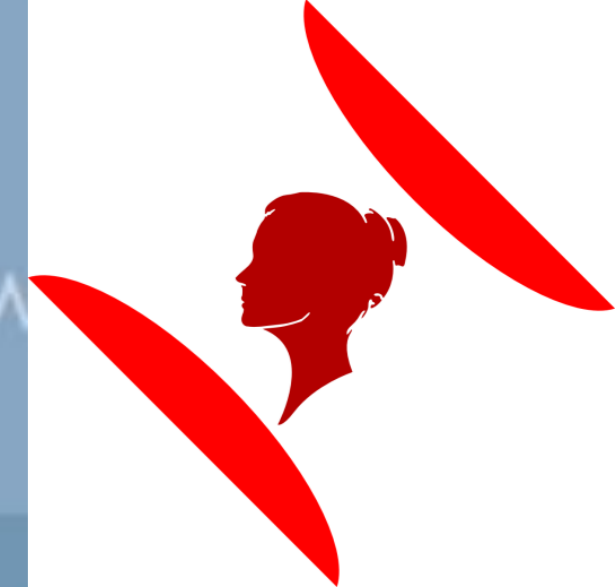


RHYTHM 2015



Arrhythmias & Heart Failure: New Insights & Technological Advances
Palais du Pharo, Marseille, France **May 28-30, 2015**

NOACs before, during and after ablation

Sok-Sithikun BUN, Decebal Gabriel Latçu, Nadir Saoudi
Princess Grace Hospital, MONACO

Congress directors

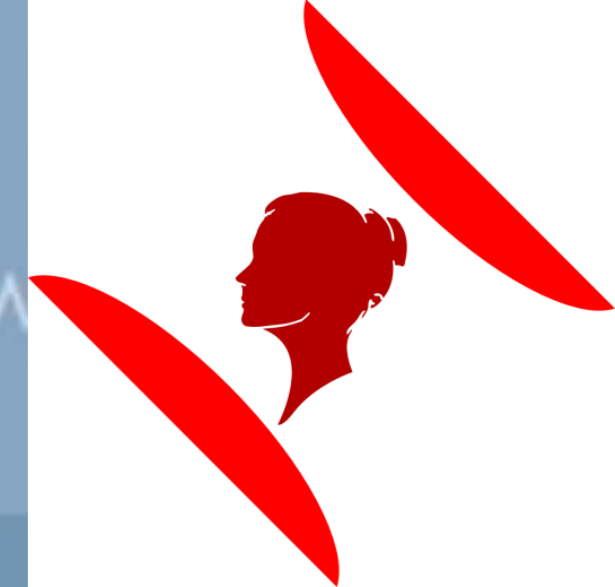
Fiorenzo Gaita
Franck Halimi
Jean-François Leclercq
André Pisapia
Julien Seitz
Jérôme Taieb

Honorary directors

Patrick Attuel
Claude Barnay



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Conflicts of interest:

Consultant fees for Daiichi Sankyo / Bayer

Congress directors

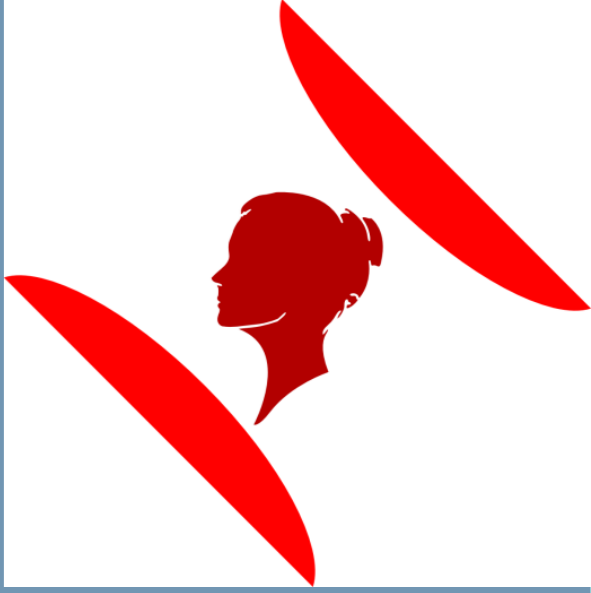
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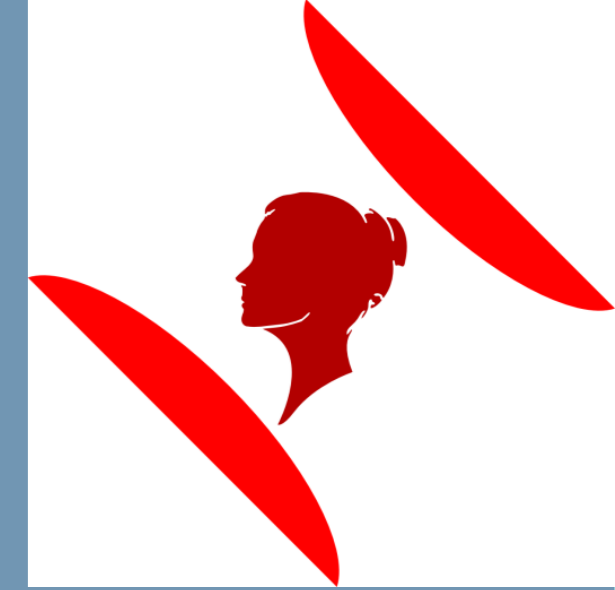


Introduction

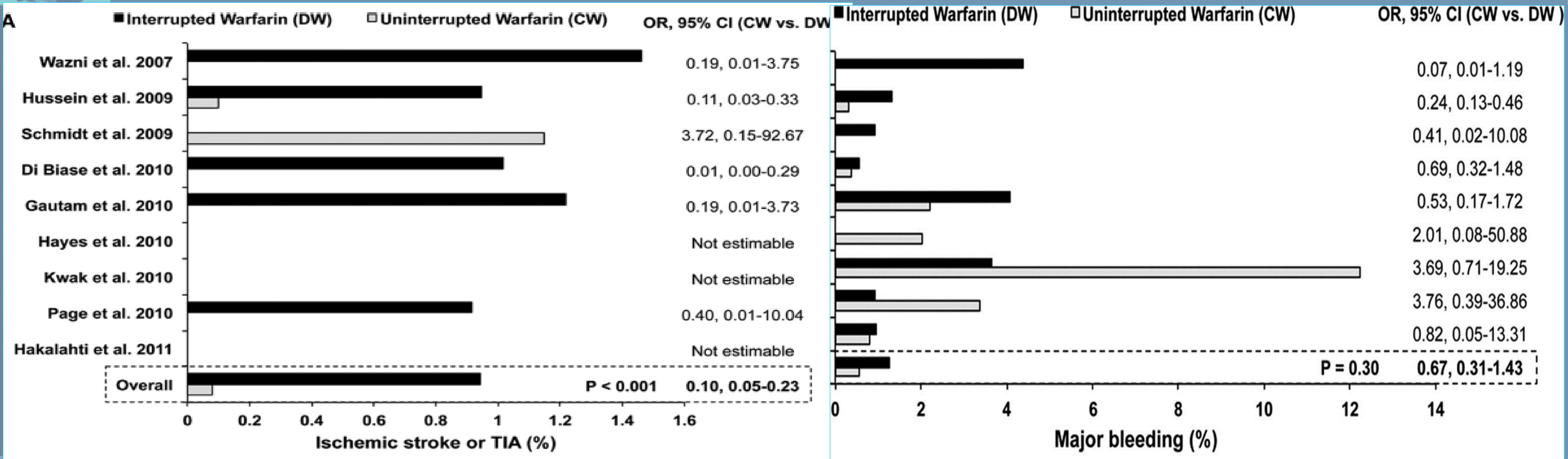


- Guidelines ?
- Feasibility / Safety / Advantages ?
- Modalities: Uninterruption ? When to resume ?
- How to manage complications ?

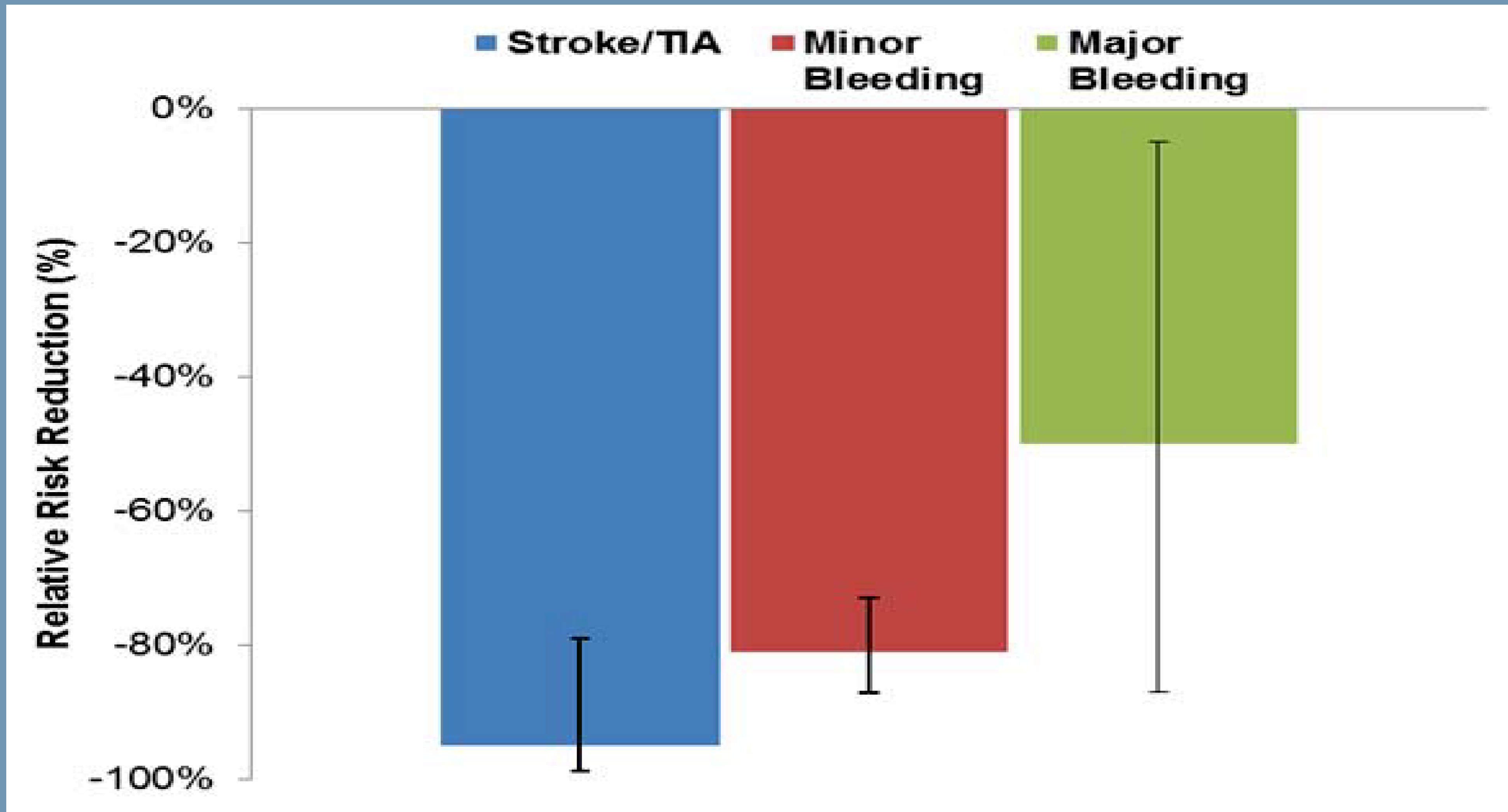
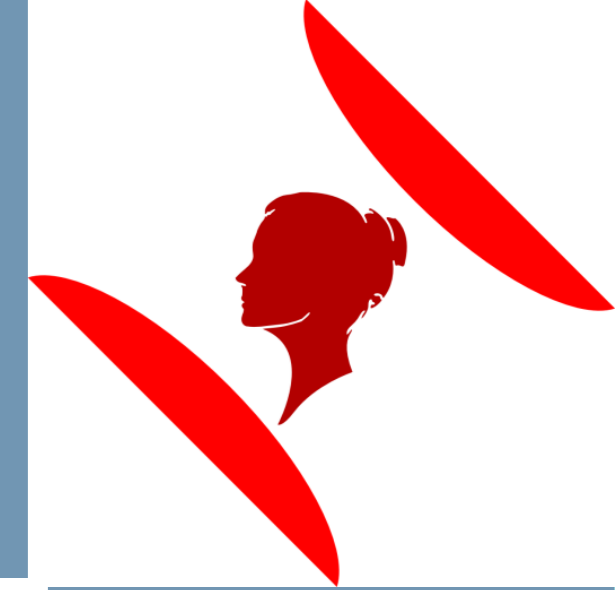
Ablation of Atrial Fibrillation Under Therapeutic Warfarin Reduces Periprocedural Complications: Evidence From a Meta-Analysis



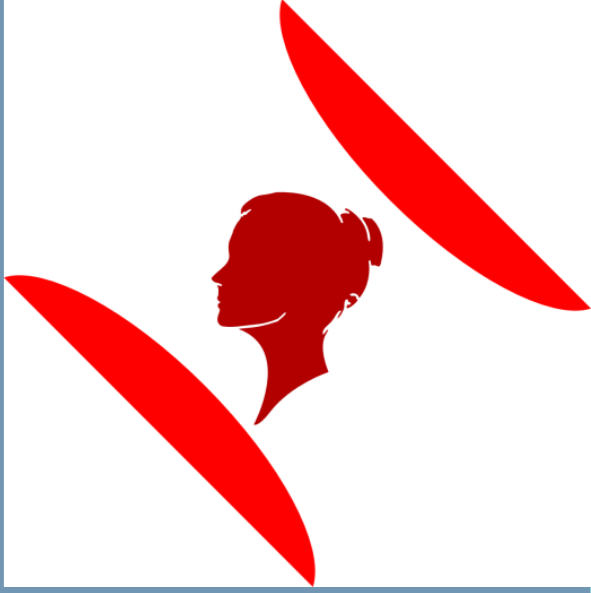
2015



COMPARE Study

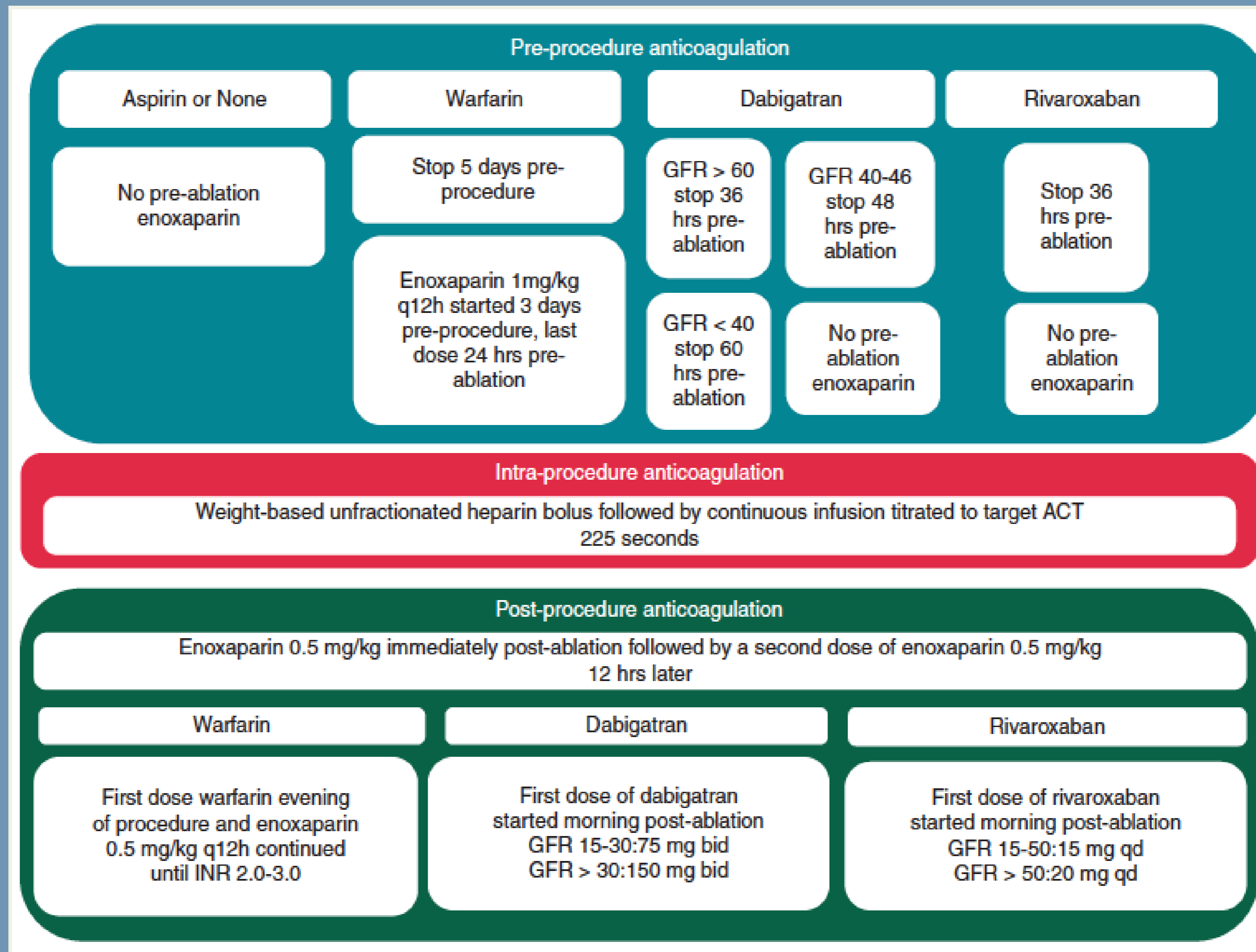
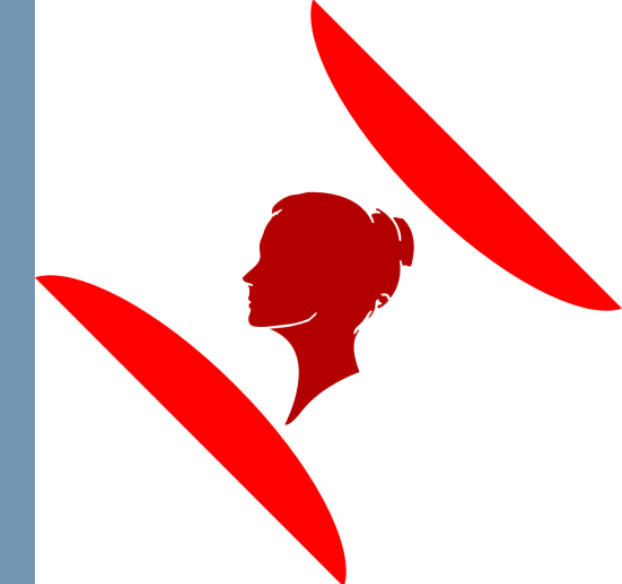


CHA₂DS₂-VASc = 0



- 214 patients
- LMWH 10 days before and 10 days after
- Long-term Aspirin
- TEE (3 %)
- 1.4 % vascular complications / No TE event

Aspirin / CHA₂DS₂-VASc = 0



EHRA practical guide on the use of NOACs in patients with non valvular AF

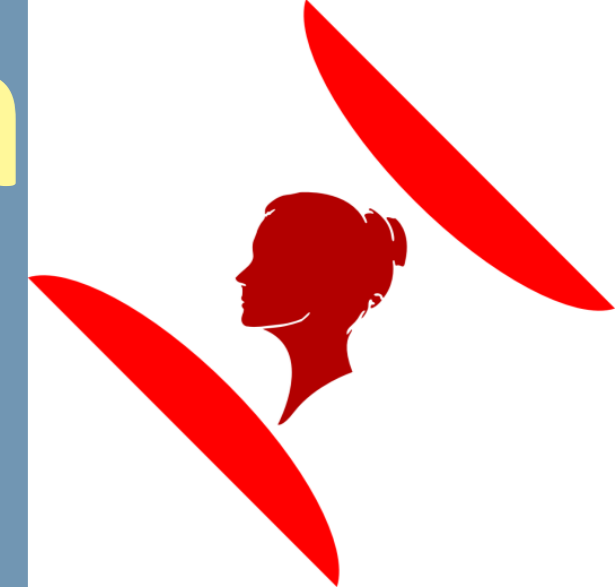


Table 9 Last intake of drug before elective surgical intervention

	Dabigatran		Apixaban		Edoxaban*		Rivaroxaban	
	No important bleeding risk and/or adequate local haemostasis possible: perform at trough level (i.e. ≥ 12 h or 24 h after last intake)							
	Low risk	High risk	Low risk	High risk	Low risk	High risk	Low risk	High risk
CrCl ≥ 80 ml/min	≥ 24 h	≥ 48 h	≥ 24 h	≥ 48 h	No data	No data	≥ 24 h	≥ 48 h
CrCl 50–80 ml/min	≥ 36 h	≥ 72 h	≥ 24 h	≥ 48 h	No data	No data	≥ 24 h	≥ 48 h
CrCl 30–50 ml/min ^b	≥ 48 h	≥ 96 h	≥ 24 h	≥ 48 h	No data	No data	≥ 24 h	≥ 48 h
CrCl 15–30 ml/min ^b	Not indicated	Not indicated	≥ 36 h	≥ 48 h	No data	No data	≥ 36 h	≥ 48 h
CrCl < 15 ml/min	No official indication for use							

Is AF ablation associated with a risk of

« major bleeding ? »

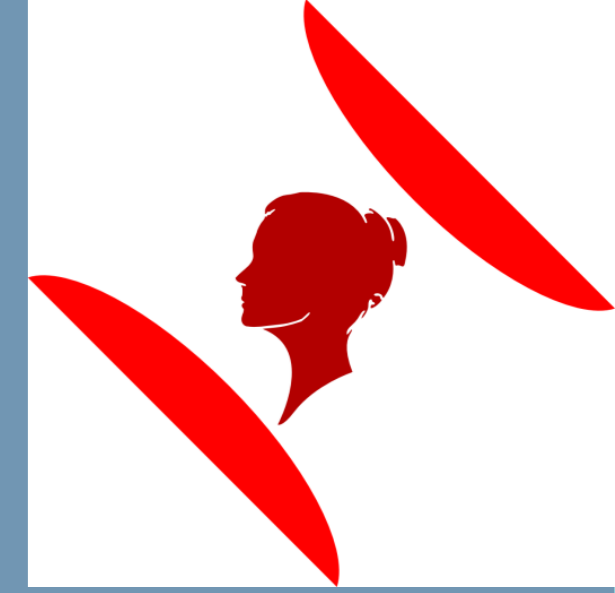


Table 10 Classification of elective surgical interventions according to bleeding risk

Interventions not necessarily requiring discontinuation of anticoagulation

Dental interventions

Extraction of 1 to 3 teeth

Parodontal surgery

Incision of abscess

Implant positioning

Ophthalmology

Cataract or glaucoma intervention

Endoscopy without surgery

Superficial surgery (e.g. abscess incision; small dermatologic excisions; . . .)

Interventions with low bleeding risk

Endoscopy with biopsy

Prostate or bladder biopsy

Electrophysiological study or radiofrequency catheter ablation for supraventricular tachycardia (including left-sided ablation via single transeptal puncture)

Angiography

Pacemaker or ICD implantation (unless complex anatomical setting, e.g. congenital heart disease)

Interventions with high bleeding risk

Complex left-sided ablation (pulmonary vein isolation; VT ablation)

Spinal or epidural anaesthesia; lumbar diagnostic puncture

Thoracic surgery

Abdominal surgery

Major orthopedic surgery

Liver biopsy

Transurethral prostate resection

Kidney biopsy

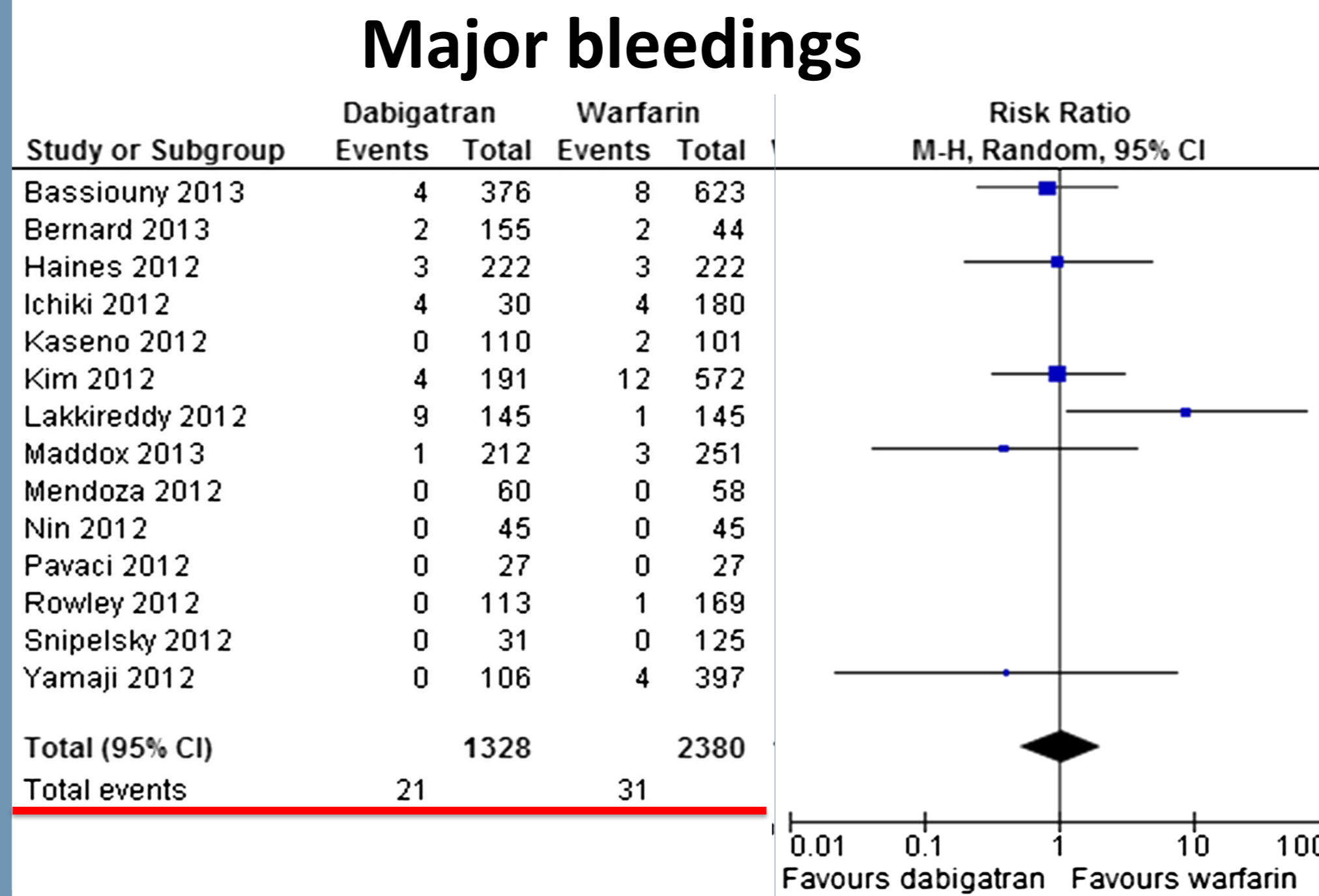
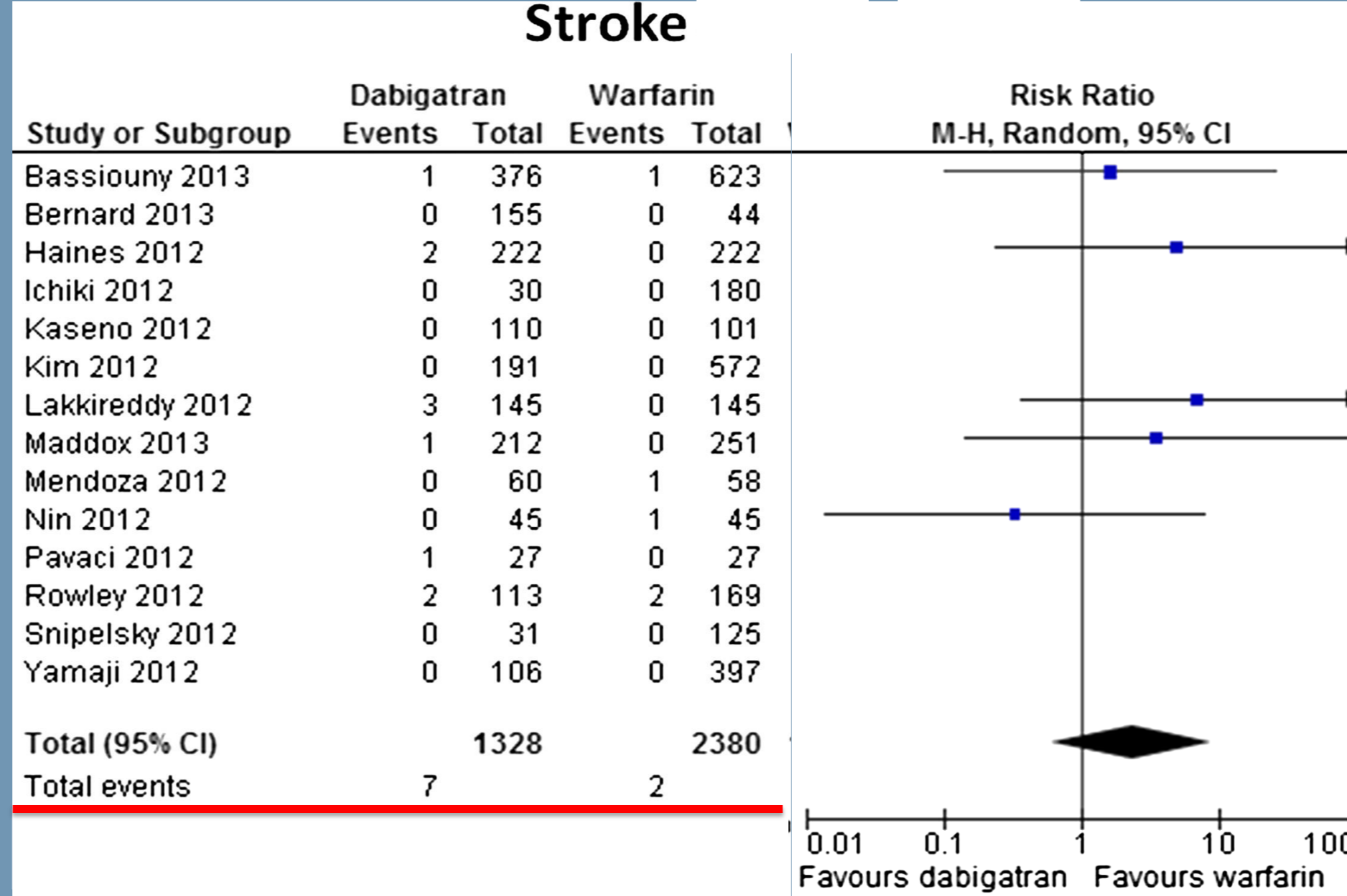
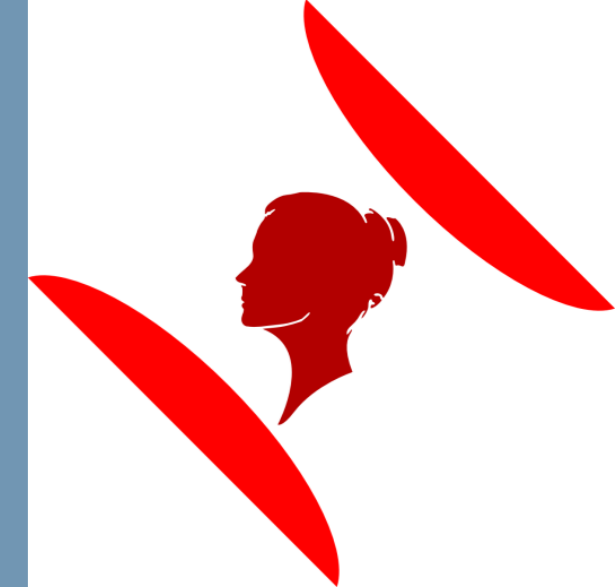
For procedures 'with a minor bleeding risk' (of which some have been listed in *Table 10*), it is recommended to discontinue NOACs 24 h before the elective procedure in patients with a normal kidney function (*Table 9*). In case of procedures that carry a 'risk for major bleeding',⁶⁶ it is recommended to take the last NOAC 48 h before.

Should catheter atrial fibrillation ablation be considered as a « high risk » intervention ?

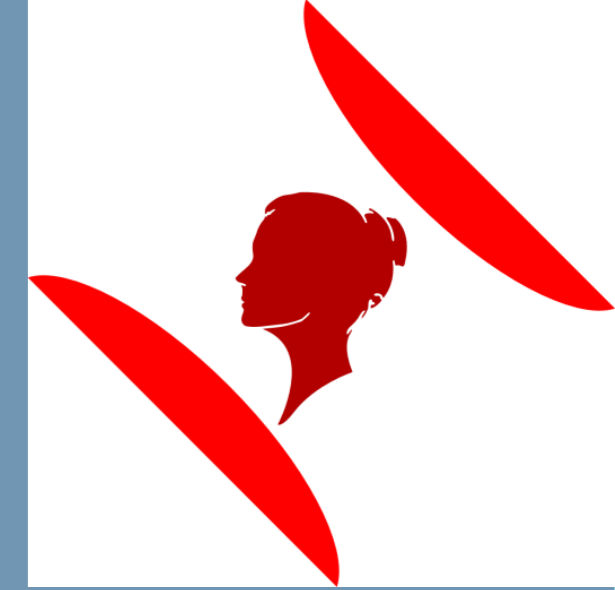
Bun SS, et al. *Europace* 2014;16(1): 150

For each patient, individual factors relating to bleeding and thrombo-embolic risk need to be taken into account, and be discussed with the intervening physician.

Dabigatran vs VKA



Rivaroxaban vs VKA

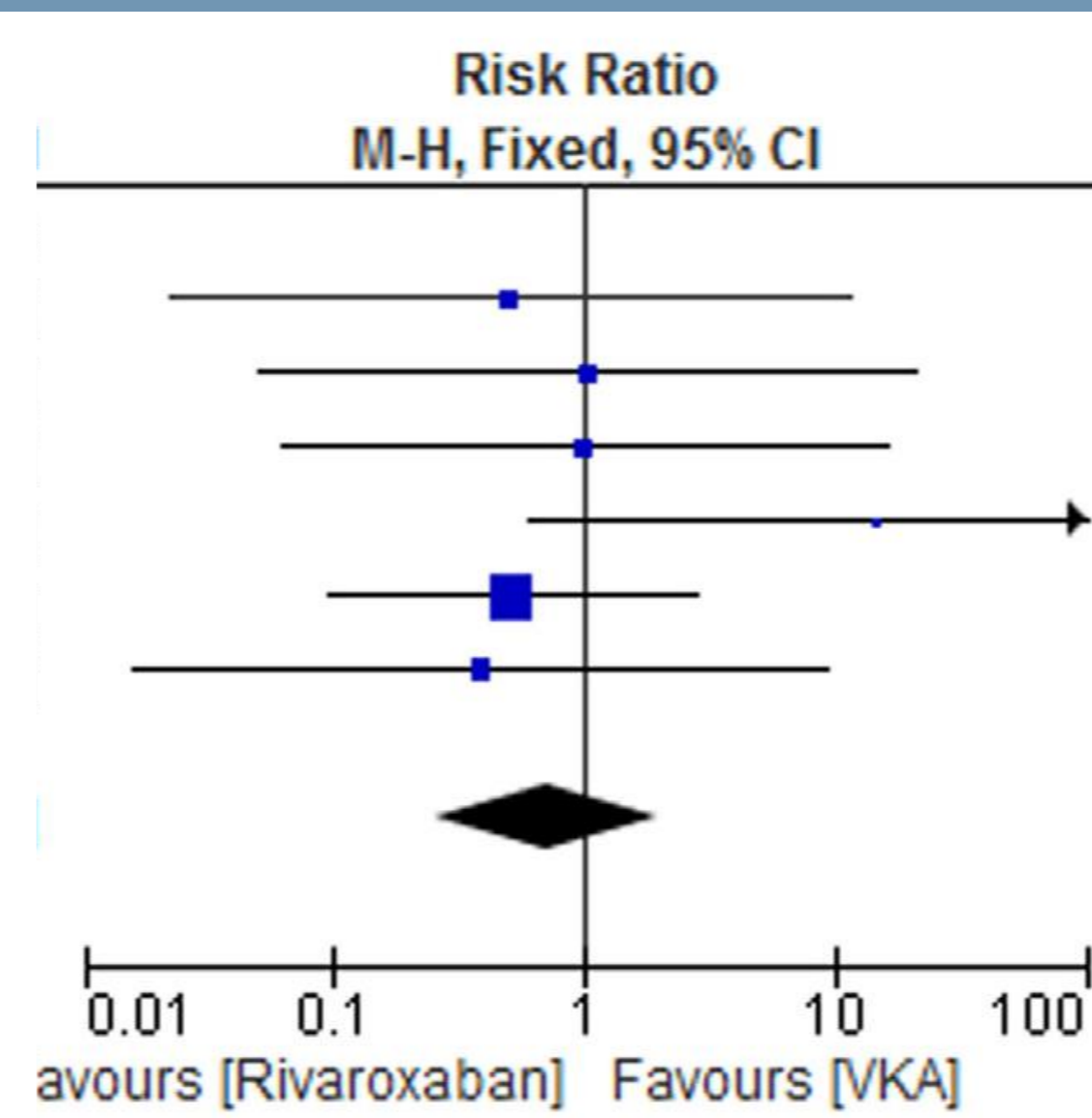


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Embolic events

A

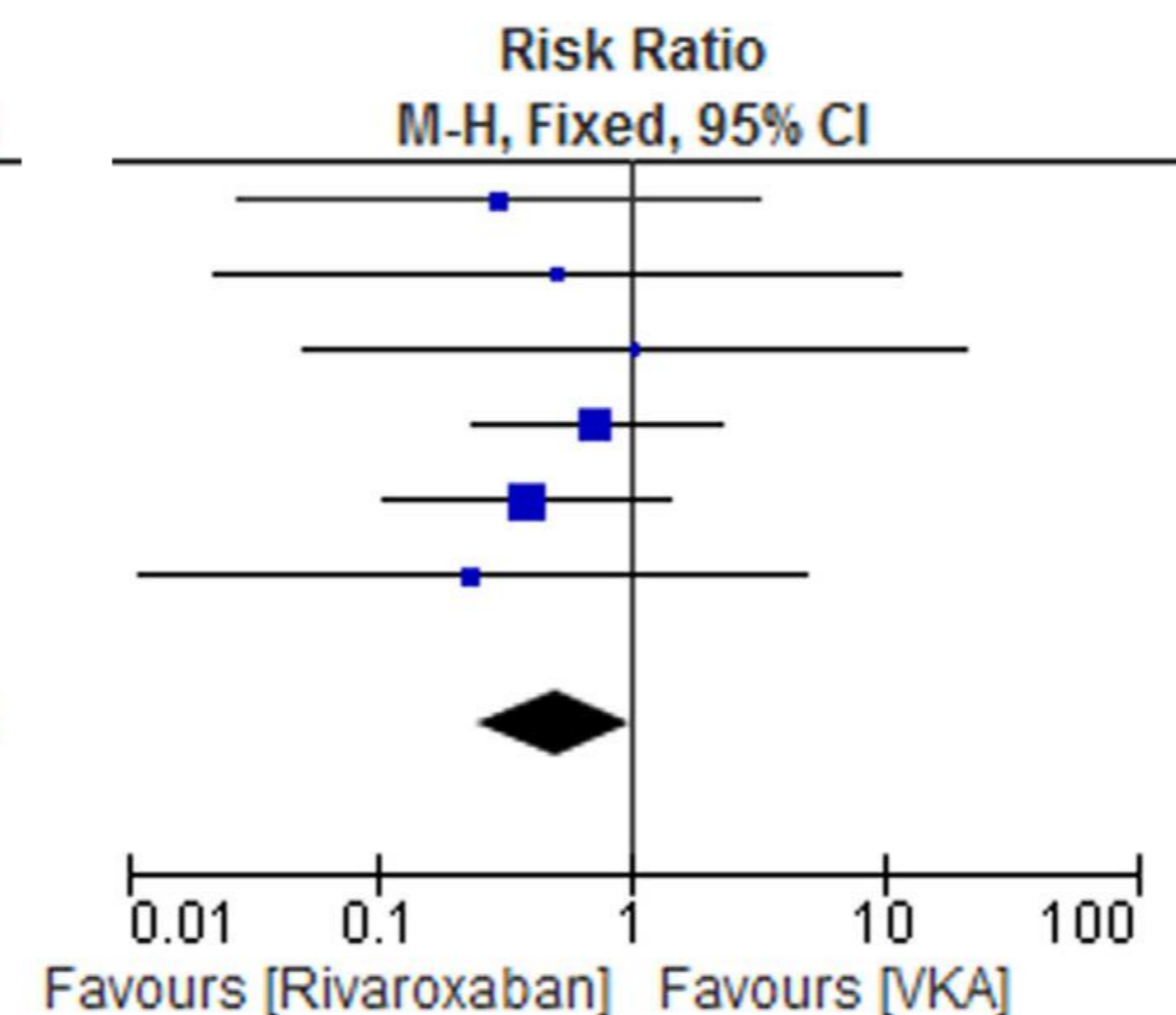
Study or Subgroup	Rivaroxaban		VKA	
	Events	Total	Events	Total
Bernard 2013	0	75	0	44
Eitel 2013	0	13	1	20
Gadiyaram 2013	0	54	2	284
Lakkireddy 2014	1	321	1	321
Murakawa 2013	0	40	1	1811
Providencia 2014	2	188	4	192
Stepanyan 2014	0	98	1	114
Total (95% CI)		789		2786
Total events	3		10	



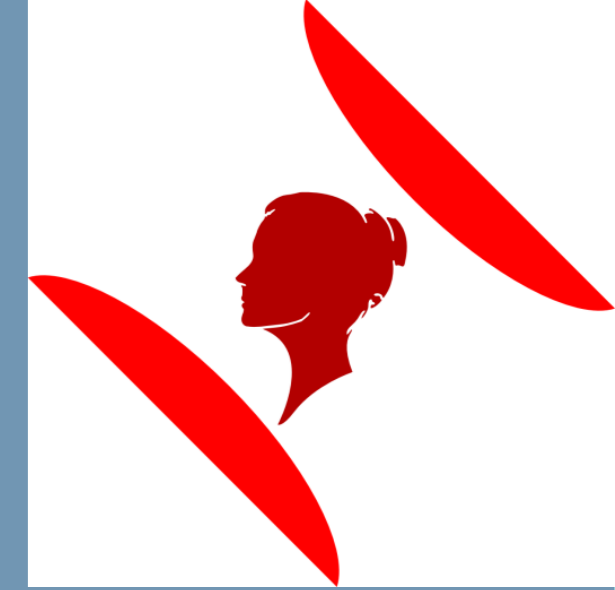
Bleeding complications

B

Study or Subgroup	Rivaroxaban		VKA	
	Events	Total	Events	Total
Bernard 2013	1	75	2	44
Eitel 2013	0	13	1	20
Gadiyaram 2013	0	54	2	284
Lakkireddy 2014	5	321	7	321
Providencia 2014	3	188	8	192
Stepanyan 2014	0	98	2	114
Total (95% CI)		749		975
Total events	9		22	



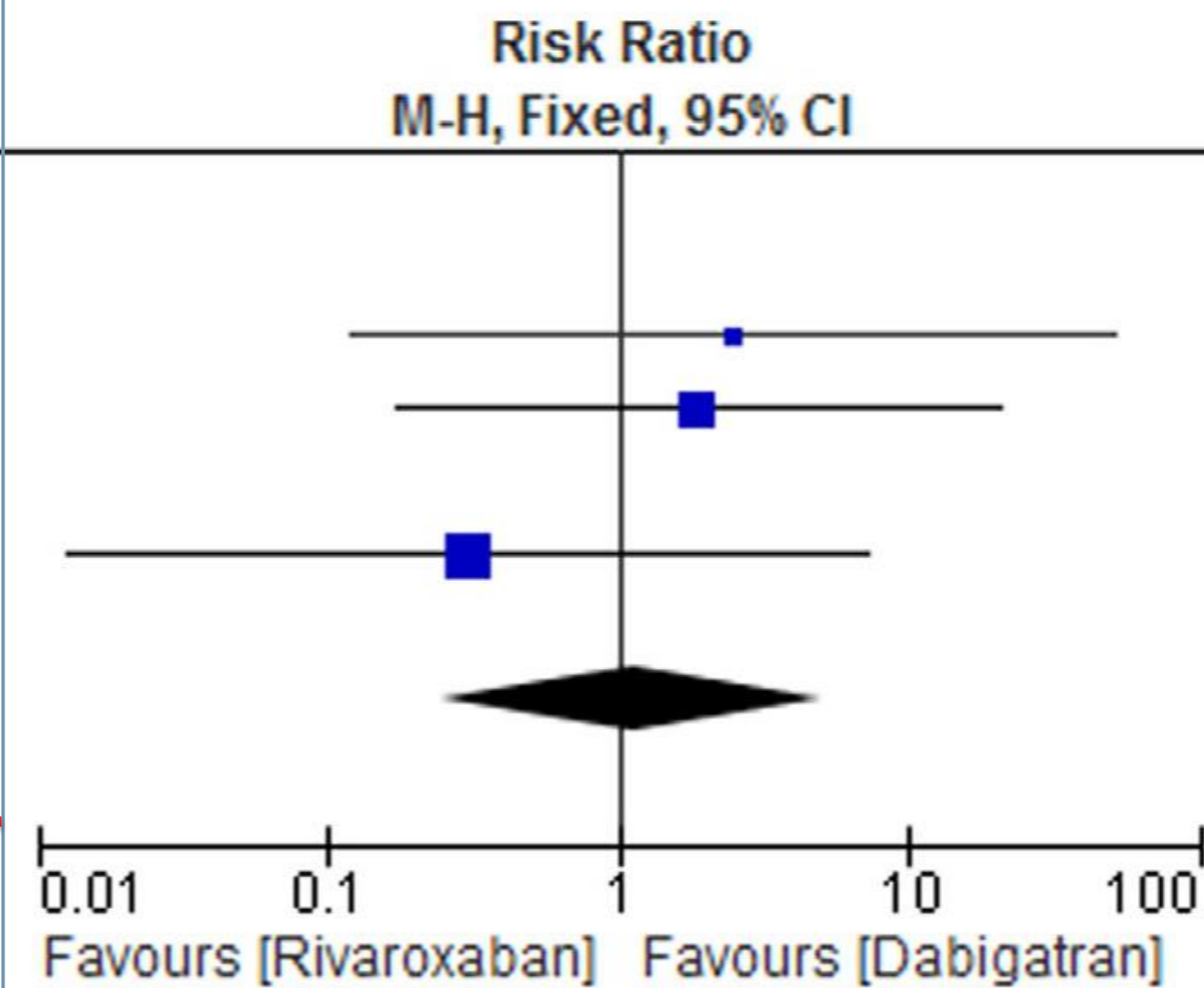
Rivaroxaban vs Dabigatran



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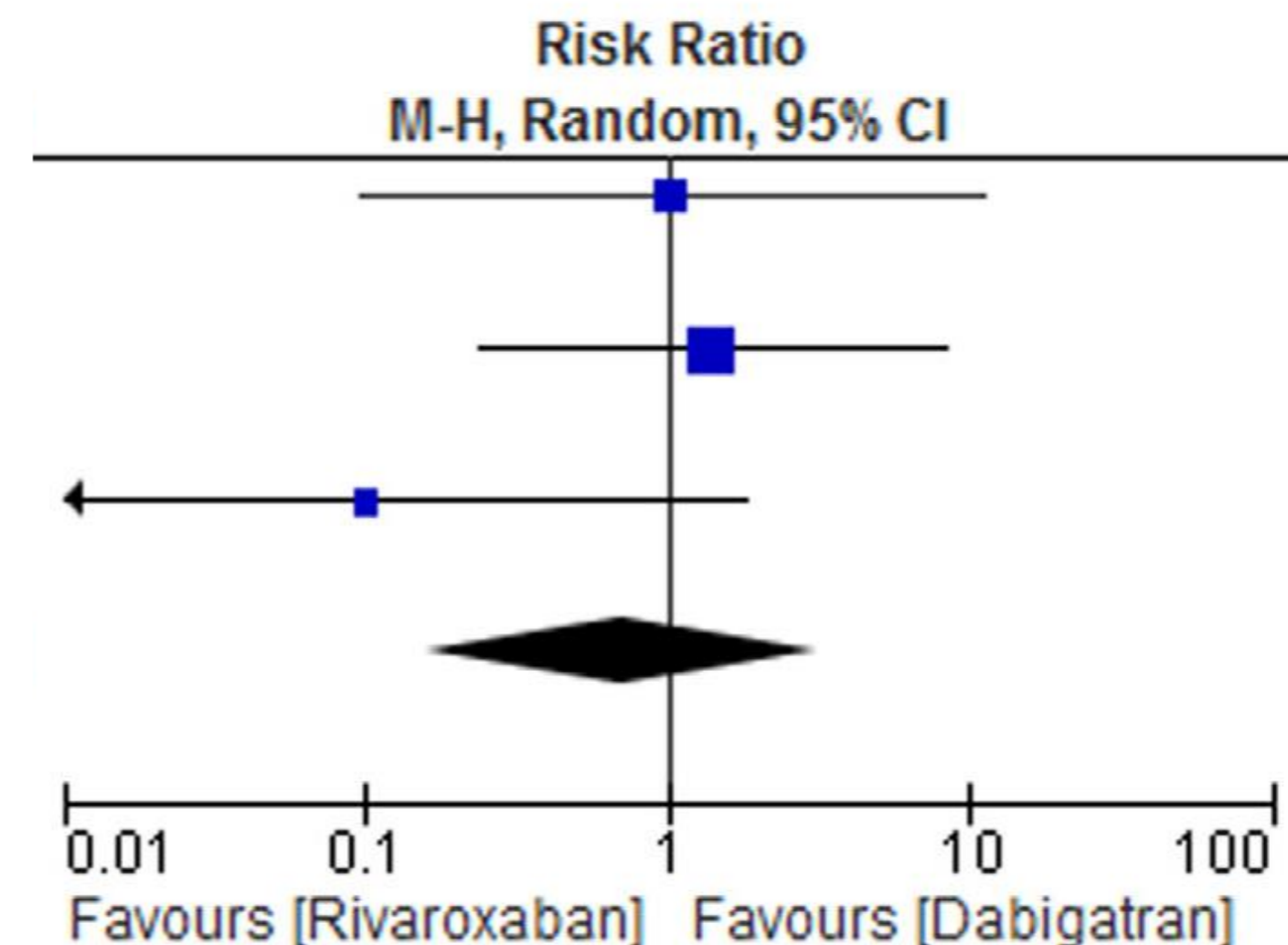
Embolic events

Study or Subgroup	Rivaroxaban		Dabigatran	
	Events	Total	Events	Total
Bernard 2013	0	75	0	155
Eitel 2013	0	13	0	41
Murakawa 2013	0	40	2	504
Providencia 2014	2	188	1	176
Sairaku 2013	0	30	0	30
Stepanyan 2014	0	98	1	89
Total (95% CI)		444		995
Total events	2		4	

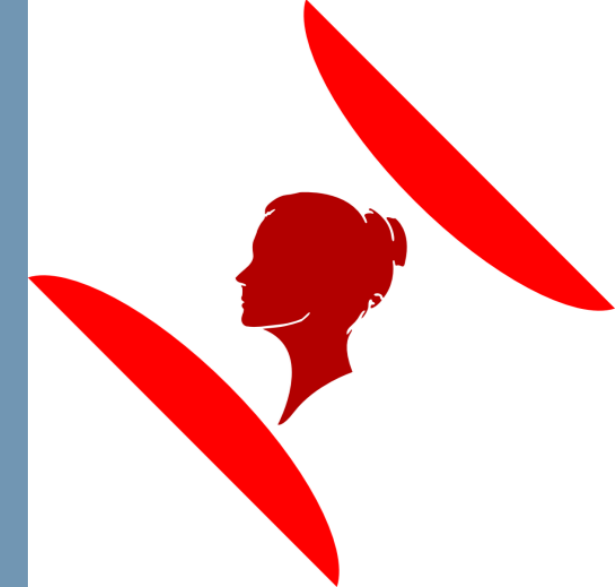


Bleeding complications

Study or Subgroup	Rivaroxaban		Dabigatran	
	Events	Total	Events	Total
Bernard 2013	1	75	2	155
Eitel 2013	0	13	0	41
Providencia 2014	3	188	2	176
Sairaku 2013	0	30	0	30
Stepanyan 2014	0	98	4	89
Total (95% CI)		404		491
Total events	4		8	

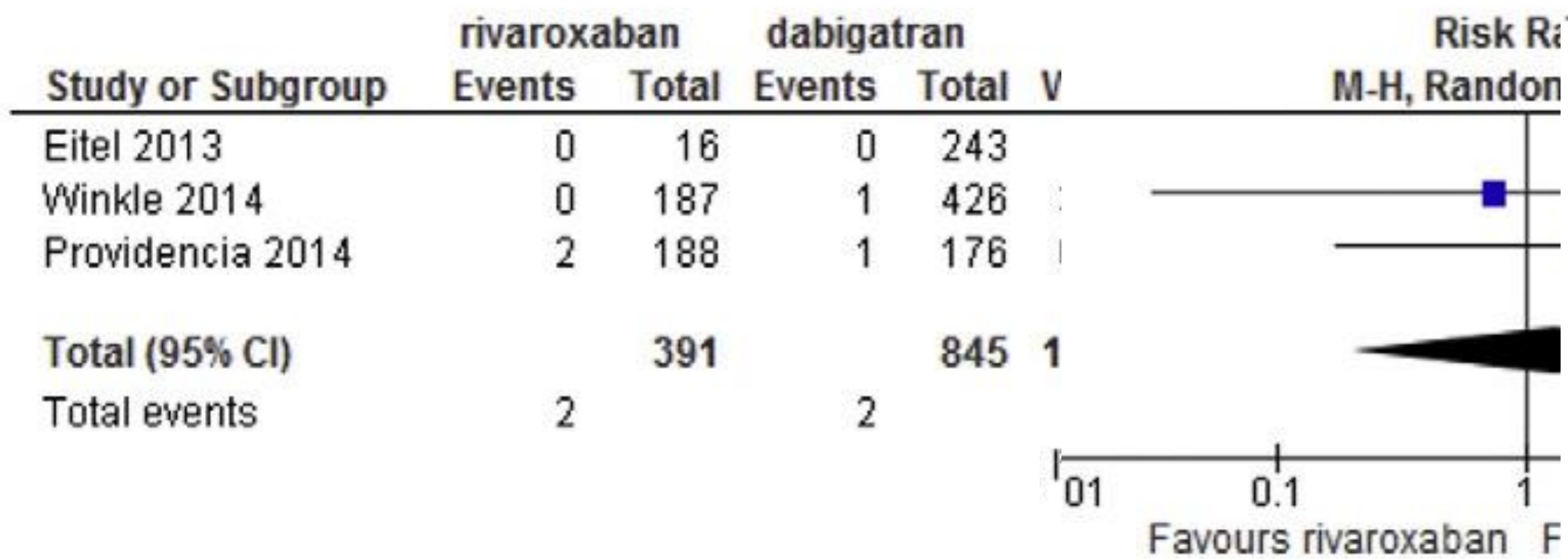
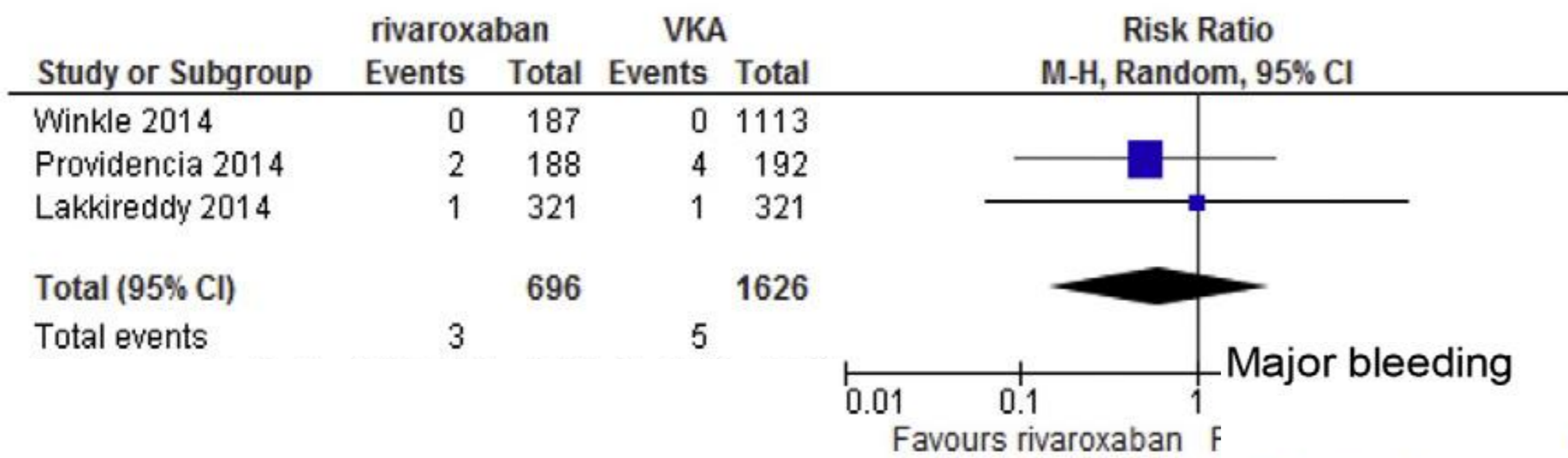
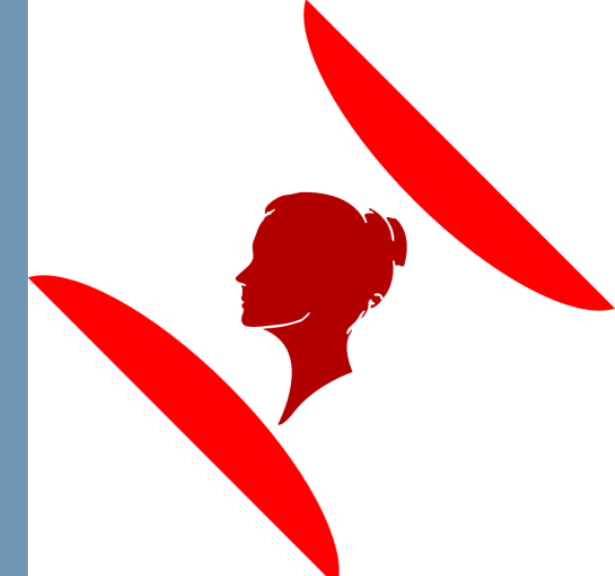


Different protocols

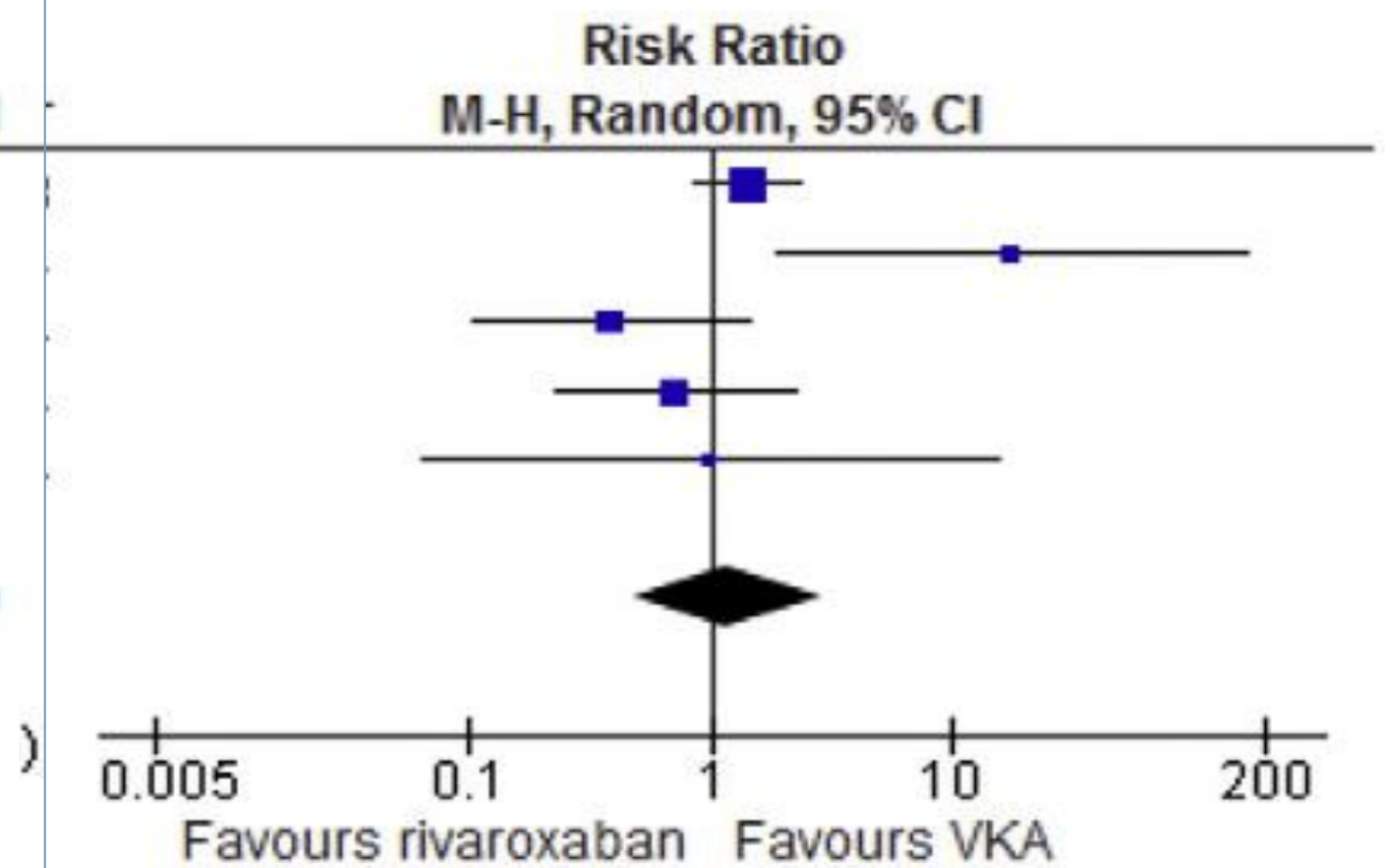


First author	Year	Study design	n (rivaroxaban)	n (VKA)	Timing of first held dose of rivaroxaban	Time interval for restarting after procedure	Target ACT (s)
Dillier	2014	R, OS	272	272	36 h prior	Morning after procedure	270–300
Winkle	2014	R, OS	187	113	36 h prior	Morning after procedure	225
Stepenyan	2014	R, OS	98	114	24 h prior	Morning after procedure	>350
Providencia	2014	P, OS	188	192	24–48 h prior	4–6 h after procedure	>300
Lakkireddy	2014	P, OS	321	321	Evening prior procedure	Evening of procedure	300–400
Piccini	2013	P, RCT	160	161	NR	NR	NR
Winkle	2014	R, OS	187	426	36 h prior	Morning after procedure	: 225
Stepenyan	2014	R, OS	98	89	2 days prior	Morning after procedure	: >350
Providencia	2014	P, OS	188	176	24–48 h prior	4–6 h after procedure	: >300
Sairaku	2013	P, RCT	30	30	24 h prior	4 h after procedure	: 300–400
Eitel	2013	P, OS	16	243	Day prior	Evening of procedure	300–350

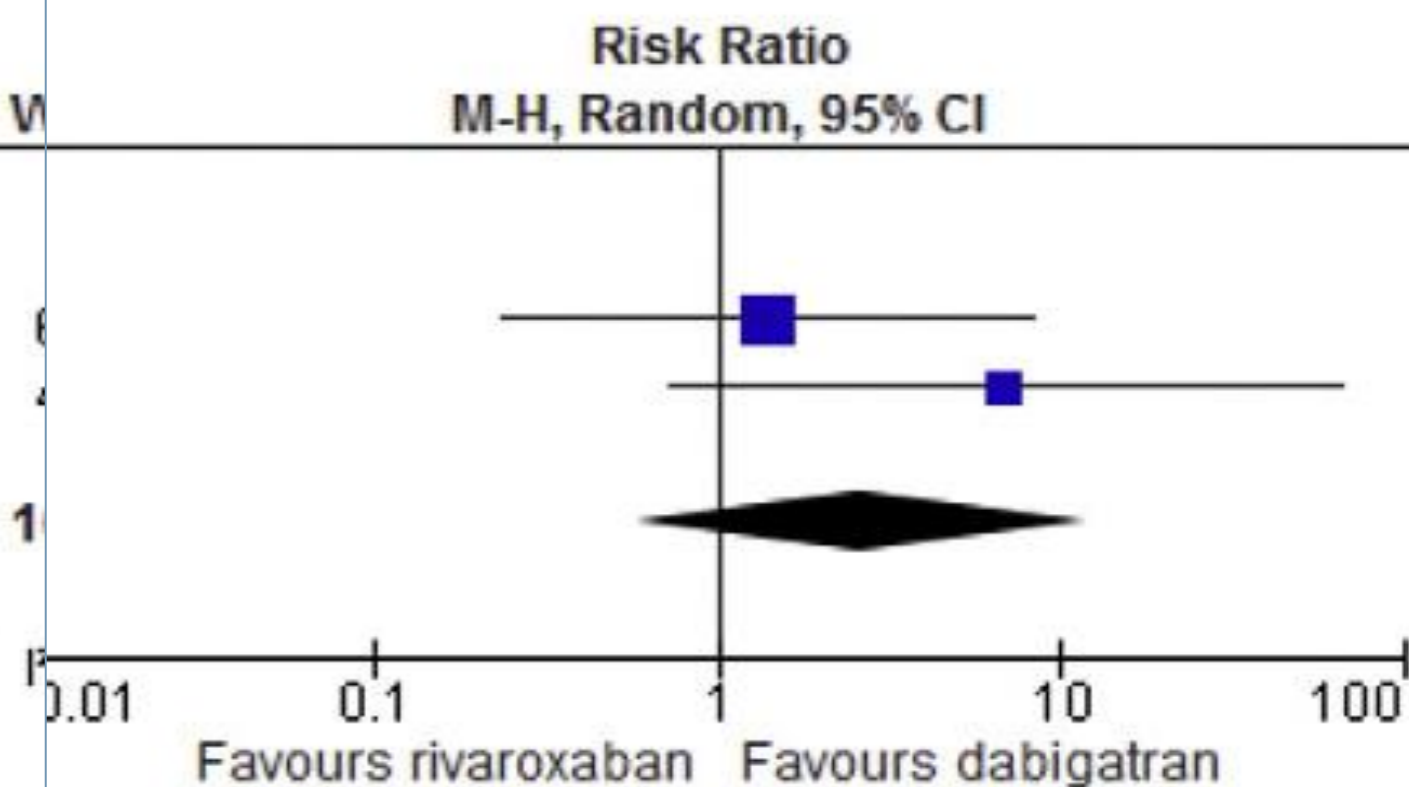
Thrombo-Embolic events



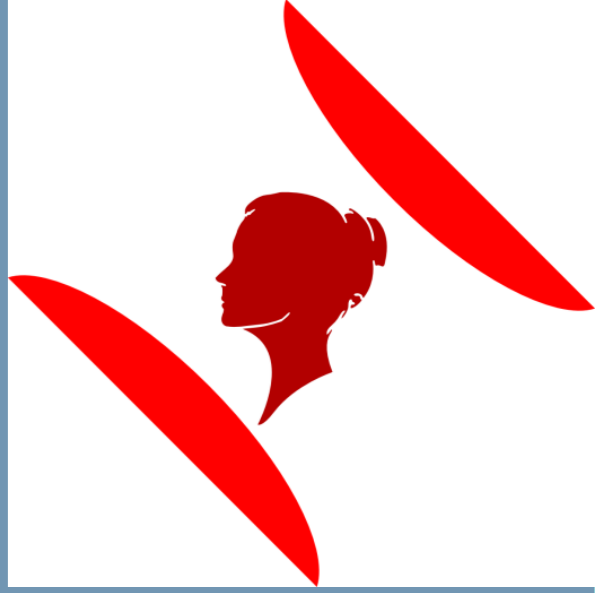
Study or Subgroup	rivaroxaban		VKA	
	Events	Total	Events	Total
Piccini 2013	30	160	21	161
Winkle 2014	3	187	1	1113
Providencia 2014	3	188	8	192
Lakkireddy 2014	5	321	7	321
Dillier 2014	1	272	1	272
Total (95% CI)		1128		2059
Total events	42		38	



Study or Subgroup	rivaroxaban		dabigatran	
	Events	Total	Events	Total
Eitel 2013	0	16	0	243
Sairaku 2013	0	30	0	30
Providencia 2014	3	188	2	176
Winkle 2014	3	187	1	426
Total (95% CI)		421		875
Total events	6		3	



Apixaban / Edoxaban



- 105 Apixaban / 210 VKA: Retrospective
 - Morning dose of 2.5 mg / Resumption in the evening
 - No difference for TE/bleeding events (10.5 vs 12.3 %)

Kaess BM et al. Am J Cardiol 2015;115:47-51.

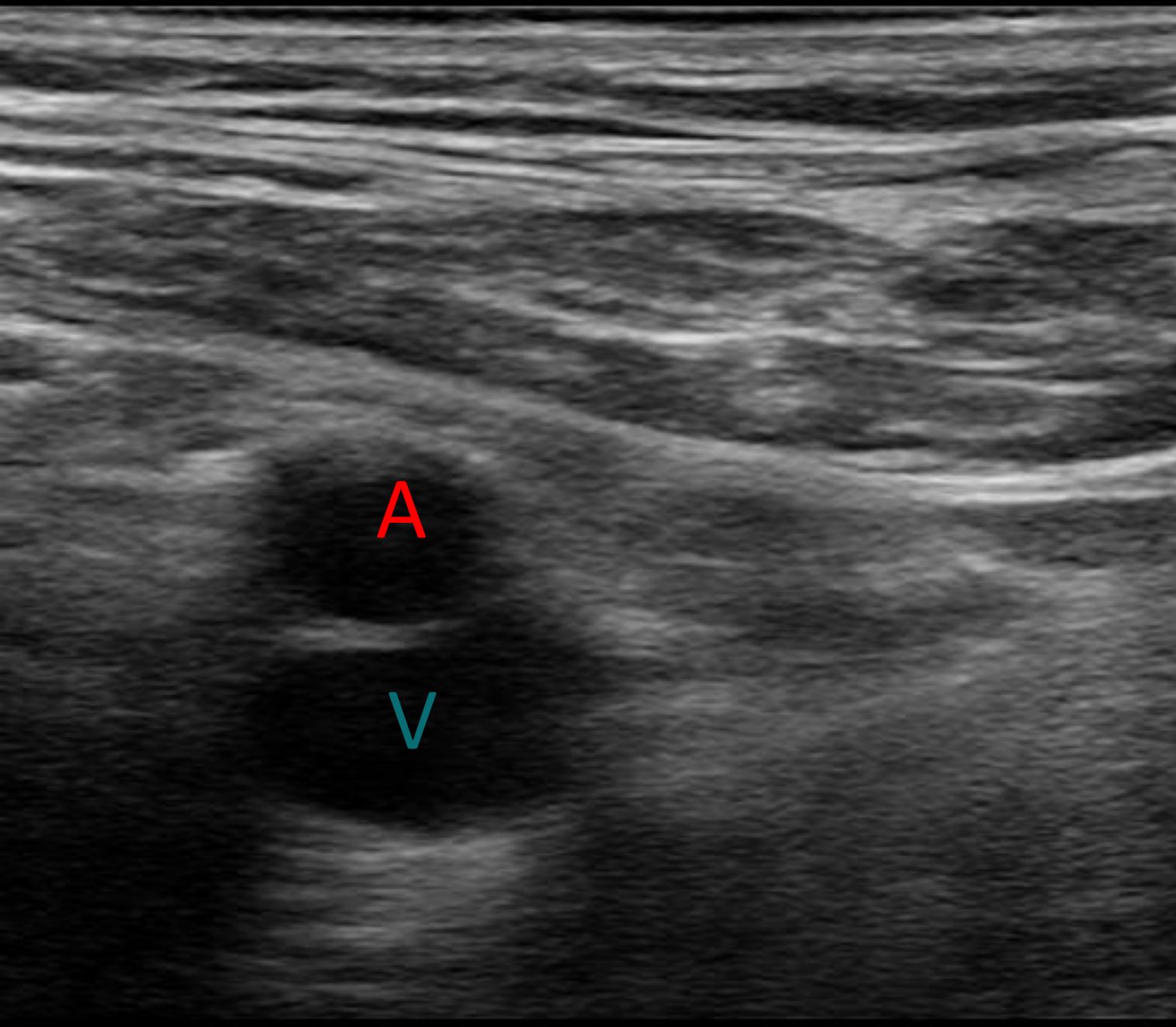
- Edoxaban: No data yet !

Ultrasound-guided venous puncture



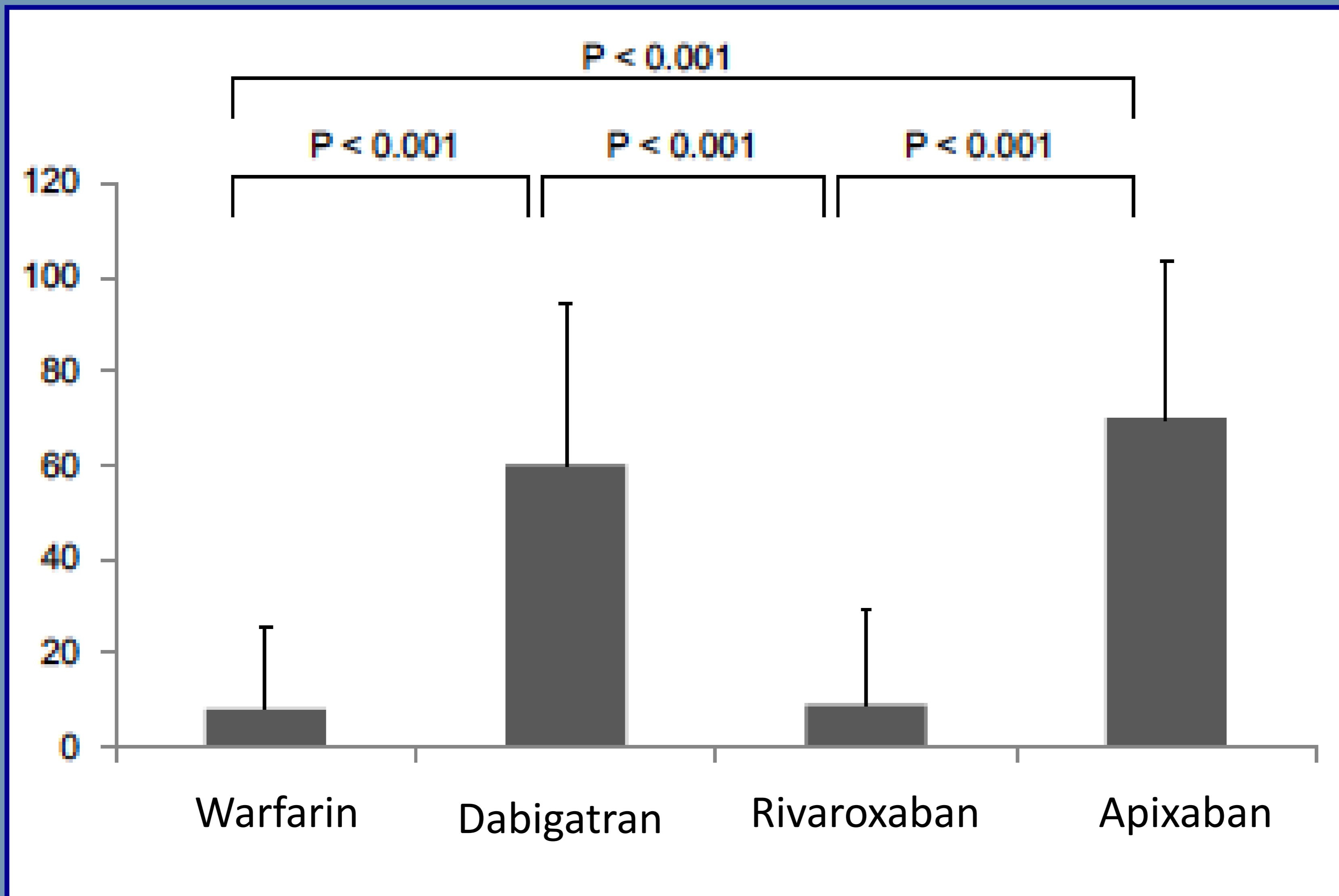
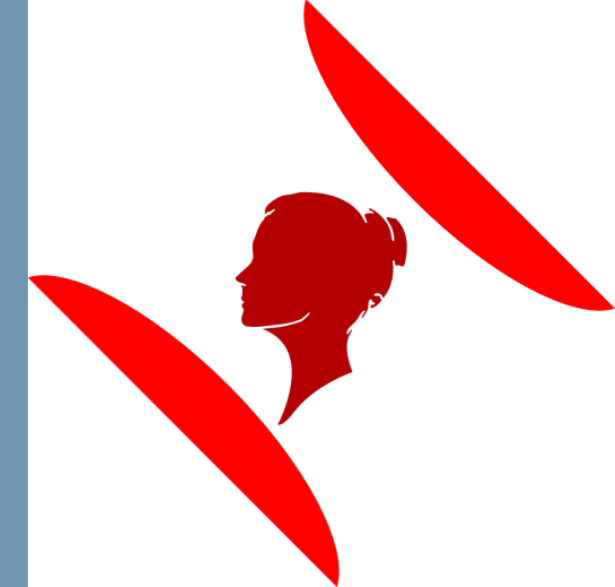
CHPG MONACO

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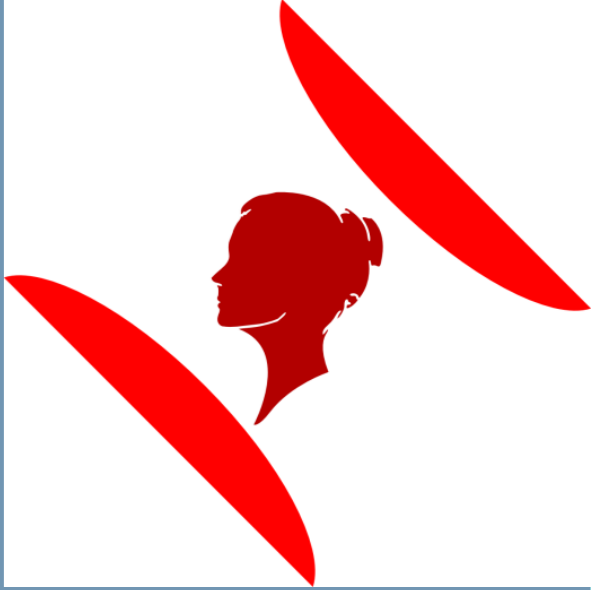
	UGVP (n=150)	No UGVP (n=150)	P
VKA (n)	82	88	
NOAC (n)	18	3	
No anticoagulation (n)	50	6	
Mean INR	2.49 ± 0.54		
Mean puncture time (s)	324 ± 145		
Mean number of sheets	3.0 ± 0.7		
Minor complications	1 minor hematoma 0.66 %	7 (4.6 %)	0.031
Major complications	0	4 (2 %)	0.042

Time to achieve ACT > 300 s



Conclusion

- Feasible and safe / Not superior to VKA
 - Several studies / Few RCTs
 - Variable protocols but same results
- When to stop NOACs before ablation ?
- Need for reversal agents ?
- Ongoing studies (RE-CIRCUIT / VENTURE-AF/AXAFA)



Welcome to the Monaco USA Arrhythmia Course 2016, March 17-19th

<http://muacmonaco.wix.com/monacousaarrhythmiaacourse>

