STUDY ELIGIBILITY FORM

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FACTORS	A	SSESSM	ENT	COMMENTS
TYPE OF STUDY				
Is the study described as observational (cohort, case control, or cross-sectional) study or randomized controled trial?	Yes	Unclear	No	
			Exclude	
PARTICIPANTS				
2. Were participants treated with warfarin?	Yes	Unclear	. No	
			Exclude	
3. Were participants of the postspecified age?	Yes	Unclear	No	Subgroups available?
NB: Please answer "Yes", If mix age participants i.e. both <=18 years and >18 years are included and state it as comments. No: If only > 18 years.			Exclude	
Genotyping				
Were study participants performed SNPs analysis (CYP2C9, VKORC1, or CYP4F2)?	Yes	Unclear	No	
			Exclude	
			LXCIUGE	
OUTCOMES				
Did the study report warfarin maintenane dose?	Yes	Unclear	No	
			Exclude	
	 			
FINAL DECISION	Include	e Unclear	Exclude	
1 X "No" = EXCLUDE				
1 X "Unclear" = UNCLEAR				

ORGANISAT	IONAL ASP	ECTS					
REF ID		Reviewer, Date Checked by					
Author, Year							
Journal/Source	e	Study ID					
Country of ori	gin						
Publication ty	ре	Fulltext / Abstract / Book chapter / internal progress report other (please specify)					
Other relevant publications in DE-form							
Fate		Decision pending / Check references / Use for discussion EX without listing / EX with listing / Other (please specify)					
Notes / Short description							
DEACONG F	OD EVOLUE	NON OF STUDY FROM		CDECIEV			
protocol	OR EXCLUS	SION OF STUDY FROM	WI REVIEW (PLEASE	SPECIFY ac	cording to		
Methods	No original	reports / Oth	ner				
Patients	No warfarin exposure age Subgroups available?						
No PGx data avilable							
Outcomes No data for relevant subgroup extractable							
Other	Duplicate publication (ID) / Other:						
NONE	Included						
OUDDENT O	TATUO.						
CURRENT STATUS Question to author							
Status verified with study investigators or sponsors: Yes							
Contact address:							

STUDY CHARACTERISTICS							
Sample size	Randomized / Recru	ited / Number ()				
Number of excluded patients	No / NA / Yes	(N=)					
Recruitment method	consecutive inclusion	/ N/A / Others	()				
Setting	in-patient / out-pati	ent / unclear	/ NR				
Location of trial							
Dates of Recruitment	From (Day/Month/Ye) To () Day/Month/Year				
Study Design		/ Cross-sectional enter trial: international ospective	/ Others / national / # centers:				
Aim of study	Warfarin dose requirement / Time to stabel dose ADE (e.g. severe bleeding or thrombosis) Others ()						
Funding	Industry / Public / Mixed (industry supported: drug / data management / travel / salary / other unclear / NR						
Conflict of interest statement	Yes 🗌 / No 🔲 / NR [
Tested SNPs	CYP2C9	VKORC1	CYP4F2				
	Yes ☐ / No ☐	Yes ☐ / No ☐	Yes 🗌 / No 🔲				
	*2 (rs1799853)	G1639A (rs9923231)	V433M (rs2108622)				
	*3 (rs1057910) Others ()	C1173T (rs9934438) Others (Others (
	P<0.05? Yes	P<0.05? Yes	P<0.05 Yes				
Definition of warfarin mainteinance dose							
Comments							

PATIENTS CHARACTERISTICS							
Age (year)	Mean () / Median () / Mode () SD () / SE () / Minimum () / Maximum () 1st Quartile () / 3rd Quartile () 95% CI inferior () / 95% CI superior ()						
Gender (Number)	Male () / Female ()						
Race (Number)	Caucasian () / Asian () African () / Others () Comment:						
Concomitant drug information	NA / Available						
Diet information	NA / Available						
Indication for warfarin	Congenital or aquired heart disease (N=) Fontan (N=) / Heart valve replacement (N=) Cardiomyopathy (N=) / Coronary aneurysms (N=) Pulmonary hypertension (N=) Other cardiac disorders (N= ,) Non-Cardiac disorder (N=) Deep vein thrombosis (N=) / Stroke (N=) Other conditions (N= ,)						
Target INR range	NA / Available						
Comments							

OUTCOME MEASUREMENT (Warfarin Mainteinance dose)										
	Data avilable	Sample number	Unit	Mean	Median	SD	MIN	MAX	95% CI Inferior	95% CI Superior
CYP2C9										
*1/*1										
*1/*2										
*1/*3										
*2/*3										
*2/*2										
*3/*3										
Variant Hetrozygus										
Variant Homozyguns										
Variant (any *2 or*3)										
VKORC1		<u>'</u>		'	1	•		•	•	•
1639-GG										
1639-GA										
1639-AA										
1639-GA or -GG										
1173-CC										
1173-CT										
1173-TT										
1173-CT or -TT										
CYP4F2	l.			<u> </u>					1	
V433M, TT										
V433M, CT										
V433M, CC										
V433M, CT or CC										

Newcastle-Ottawa Quality Assessment Scale for Cohort Studies

Selection

- 1) Representativeness of the variant genotypes in each SNP
 - a) truly representative in the community *
 - b) somewhat representative in the community *
 - c) selected group of users
 - d) no description of the derivation of the cohort
- 2) Selection of the wild genotype in each SNP
 - a) drawn from the same community as the exposed cohort *
 - b) drawn from a different source
 - c) no description of the derivation of the non exposed cohort
- 3) Ascertainment of genotype
 - a) secure record ★
 - b) structured interview ★
 - c) written self report
 - d) no description
- 4) Demonstration of genotyping was not present at start of study
 - a) yes ☆
 - b) no

Comparability

- 1) Comparability of cohorts on the basis of the design or analysis
 - a) study controls for body size or age ★
 - b) study controls for any additional factor (e.g. target INR range, indication, or diet) *

Outcome

- 1) Assessment of outcome
 - a) independent blind assessment *
 - b) record linkage ☆
 - c) self report
 - d) no description
- 2) Was follow-up long enough for outcomes to occur
 - a) yes (more than 1 month of stable warfarin control) 🖈
 - b) no
- 3) Adequacy of follow up of cohorts
 - a) complete follow up all subjects accounted for at least 1 months ★
 - b) subjects lost to follow up unlikely to introduce bias small number lost <10 % ★
 - c) follow up rate < 90% and no description of those lost
 - d) no statement

Total number of stars: stars