## PQS Oral Anticoagulant Safety Audit 2023/24 - Data Collection Form

Sect	ion 1 - All patients	
1.	Patient's name (For internal use – not for reporting to NHS England)	
2.	Date	/ /
3.	Patient's age	
4.	Is the patient a care home resident?	Yes No Not known
5.	Name of anticoagulant that the patient is taking*	Acenocoumarol Phenindione
		Apixaban Rivaroxaban
		☐ Dabigatran ☐ Warfarin
	*for patients prescribed more than one anticoagulant, see question 7. This does not include where a patient is prescribed two strengths of the same medicines to make a dose e.g., multiple strengths of warfarin	Edoxaban
6.	Is the anticoagulant supplied in a monitored dosage system / compliance aid?	No
		Yes, one medicine per blister / compartment
		Yes, multiple medicines per blister / compartment
7.	Is the patient prescribed more than one ora	I No (go to question 8)
	anticoagulant? (Please do not include a patient prescribed two strengths of the	Yes
	same medicine to make a dose e.g.,	Name of other anticoagulant:
	multiple strengths of warfarin)	What action did you take and what was the outcome?
		If patients are switching anticoagulant treatments, remind them to return any medicine no longer needed for safe disposal.
	Is the patient prescribed an oral NSAID* as	☐ No (go to question 9)
	* <b>Do not</b> include low dose aspirin (300mg or less per day) here; record it in Q10 instead.	☐ Yes ☐
		8b. Is the patient also prescribed gastro-protection?  (e.g. a proton pump inhibitor or H2 receptor antagonist)
		Yes
		□ No
		<b>8c.</b> Have you contacted the prescriber about concomitant use of an anticoagulant with an NSAID?
		☐ Yes – prescriber discontinued anticoagulant and/or NSAID
		Yes – prescriber confirmed no medication changes required
		Yes – gastro-protection prescribed
		Yes – other action by prescriber. Please specify:
		□ No – please specify the reason:

9а.	well as the anticoagulant?		☐ Yes ☐		
			(e.g. a proton pump into The PINCER summary	also prescribed gastro-protection? nibitor or H2 receptor antagonist) nio indicates that gastro-protection should and offered when combination therapy iplatelet) is indicated.	
			☐ Yes ☐ No ☐  9c. Have you conta	acted the prescriber for a review of	
			gastro-protection?		
				rotection prescribed er discontinued anticoagulant and / or	
				er confirmed no medication changes	
			-	r has been contacted about gastro- nis patient within the last 6 months	
				is discussed with prescriber and has not to take gastro-protection	
				ason. Please specify:	
			No – other reas	son. Please specify:	
10.	Which category best describes how the audit	_	ion with the he pharmacy	∑ Go to	
	was completed for this patient?		on with the	Section 2	
	patient	patient by t	telephone ion with the		
		patient by			
			th patient by e, e.g. email		
		Patient's repharmacy, contact par		VKA prescribed – Go to Section 3  DOAC prescribed – Go to Section 4	
		Medicine d pharmacy, contact pa			
		to contact	e patient, unable patient / tive / care staff		

Section 2 - Patient feedback (only complete this section if you can contact the patient)						
11.	Was the patient already awa taking an anticoagulant, i.e. a the blood/prevent blood clots	a medicine to thin		res No – information provided No – information not provided		
12.	Did the patient already know over-anticoagulation, e.g. un bruising, nose bleeds?			res No – information provided No – information not provided		
13.	Was the patient already awa check with the doctor or phataking over-the-counter mediproducts or supplements?	rmacist before		Yes No – information provided No – information not provided		
14.	For patients taking vitamin K Was the patient already awa change can affect their antic medicine?	re that dietary		Yes No – information provided No – information not provided Not applicable		
15a.	anticoagulant alert card?	ticoagulant Alert Card  sharp entirespectate therepy to provide the through the transport of the transport o		Yes, card seen by pharmacy staff Yes, card not seen but patient confirmation they have this card No card but aware of card No card and unaware of card		
	Transment of a rist  Name and riskform  Transport or control  Tran	Transmittation of the property		. Was a standard yellow alert card offered to patient?		
				Yes, card accepted Yes, but card declined because the patient has manufacturer's alert card Yes, but card declined because the patient has another anticoagulant alert card Yes, but card declined for other reason No, not offered. Reason - please specify		
Vitamin K antagonist prescribed? Go to Section 3 DOAC prescribed? Go to Section 4						

Section 3 - Patients prescribed vitamin K antagonists only						
16a.	Did you find out <u>when</u> the patient last had an INR test before issuing this medicine?	☐ No (go to question 16d) ☐ Yes ☐				
16b.	How did you obtain this information? (select all that apply)	From patient From patient's representative From yellow anticoagulant record book or other written record From general practice From patient's care provider, e.g. nursing home From anticoagulant service From other source - please specify:				
16c.	How long ago was the INR test?	Fewer than 4 weeks (go to Section 4)  4 – 12 weeks (go to Section 4)  More than 12 weeks				
	If the INR test was more than 12 weeks ago, what, if any, action did you take?	(go to Section 4)				
Section 4 – All patients						
17.	Please give details of any other referrals or action taken about anticoagulant safety issues, e.g. drug interactions, INR concern (do not include any patient identifiable information)					