

APPENDIX D – Annual Review Report Form (May 2022)

Oral Anticoagulation Clinic Details			
Service Provider Name			
Address Of Anticoagulation Clinic			
Telephone Number		Facsimile	
E-Mail			
Name of Lead Clinician			
Name of Person Compiling Report		Position/Post	
Patient Statistics			
1.	Total Number of Patients managed in the service during past 12 months		
2.	Number of Initiations by the Service Provider / within Primary Care		
3.	Number of Referrals Received from Secondary Care		
4.	Number of Referrals referred back by the Service Provider to Secondary Care		
5.	Number of adverse incidents reported		
	Clinical outcome:		
	Death		
	Severe (permanent harm)		
	Moderate (significant, but not permanent harm, requiring increase in treatment)		
	Low (temporary harm, requiring extra observation or minor treatment)		
	No harm		
	Total		
	Type of report:		
	Prescribing		
	Dispensing/medicine preparation		
	Administration		
	Monitoring		
	Total		
	Type of incident:		
	Wrong dose		

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	Wrong frequency	
	Omitted medicine/dose	
	Wrong drug	
	Wrong quantity	
	Mismatching of patient and their medicine	
	Wrong/transposed/omitted medicine label	
	Wrong/omitted/passed expiry date	
	Wrong storage	
	Wrong route	
	Contraindication	
	Patient allergic to treatment	
	Wrong formulation	
	Wrong method of preparation/supply	
	Adverse drug reaction – when used as intended	
	Wrong or omitted verbal patient directions	
	Other	
	Total	
6.	Number of Serious Untoward Incidents reported	
7.	Number of Episodes Requiring the Administration of Vitamin K	
8.	Total Number of Domiciliary Patients	
9.	Number of Domiciliary Visits undertaken	
10.	Number of individual INR Tests Completed	
Patient Experience		
11.	Number of formal patient complaints received by Service	
12.	Number of informal patient complaints received by Service	
13.	How many patient complaints have you received during this time period which related to:	
13a.	Communication / information to patients	
13b.	Clinical treatment	
13c.	Attitude of staff	
13d.	Appointments, delay / cancellation	
13e.	Patient privacy, dignity	
13f.	Referral, discharge and transfer arrangements	
13g.	Please give an example of actions taken in response to a complaint	

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14.	Number of patients included within the Patient Satisfaction Survey within the year Of those patients, number of responses received	
15.	Any issues of low patient satisfaction (e.g. waiting times / accessibility / continuity of care)	Yes / No*
16.	If you have answered YES to question 6, provide details in the space provided below (Additional space is available on Page 7 of this document)	
<i>*questions 14-16 are not required IF the service is participating in the PROM Light CQUIN</i>		
17.	Service changes made in response to patient feedback	Yes / No*
17a.	If you have answered YES to Question provide details in the space provided below (Additional space is available on Page 7 of this document)	
Risk Assessments & Reporting Procedures		
18.	Are Clinic Risk Assessments Up-to-date?	Yes/No*
18a.	If you have answered NO to Question 12 give details of the date of last assessment and any remedial actions that are being taken in the space below (Additional space is available on Page 7 of this document)	
18b.	If you have answered YES to Question 12 enter the date of the most recent assessment	
19.	Has the Service Provider completed 2 6 monthly cycles of the audit toolkit (Appendix D)?	Yes/No*
20.	At the time of writing have <u>all</u> Adverse Incidents been investigated, lessons identified and responded to, and the incident completed satisfactorily?	Yes/No*
20a.	If you have answered NO to Question 14 give BRIEF details in the space below (Additional space is available on Page 7 of this document)	

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Equipment & Service		
21.	Which Computer Assisted Decision Support Software is used?	
21a.	Give details of the software below & any issues arising	
21b.	If you have answered YES to Question 16 give brief details about how you consider its efficiency compared to clinical decision making (Additional space is available on Page 7 of this document)	
22.	In the space below give detail of the Point Of Care Testing (POCT)equipment used in the service (Additional space is available on Page 7 of this document)	
23.	In the space below describe what quality assurance checks are carried out to ensure that the POCT equipment is functioning correctly (Additional space is available on Page 7 of this document)	
23.a	Confirm registration with NEQAS & provide a copy of the annual report or evidence of data submitted	
24.	In the space below give BRIEF details of any equipment failures and what contingency actions were taken or if failures have not occurred what contingency plans are in place (Additional space is available on Page 7 of this document)	
25.	Were any equipment failures reported to the Medicines and Healthcare Products Regulatory Agency (MHRA) during this period	Yes / No*
Training And Accreditation		

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26.	Are the training needs of all staff assessed and documented on an annual basis.	Yes/No*
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27.	As a minimum, are all staff involved in the delivery of clinical care to patients, trained in accordance with the relevant stated NPSA Workforce Competencies	Yes/No*
27a.	If you have answered NO to Question 22 give BRIEF details of Training Plans in the space below (Additional space is available on Page 7 of this document)	
28.	Have any staff members completed e-learning packages or attended any courses relating to Oral Anticoagulation Therapy.	Yes/No*
28a.	Please give details below, of courses attended and the designation of the attendee. (Details to include the full name of the course, the accrediting authority and date the course was completed.) (Additional space is available on Page 7 of this document)	
29.	Do you hold a copy of the latest NPSA Alert pertaining to Oral Anticoagulants and is this document readily available to all staff?	Yes/No*
30.	Are documented procedures for all key areas identified within paragraph 4.3.6 of the specification, held by the service provider and readily available to all staff	Yes/No*
General		

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31.	In the space below enter any other observations regarding the Oral Anticoagulation Service relevant to the Annual Review. (Additional space is available on Page 7 of this document)

(* Please delete as appropriate)

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Additional space for answers. Please annotate the question number clearly in the margin

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