Amendment Tool

v1.6 06 December 2021

Short project title*:										
IRAS project ID* (or REC reference if no IRAS project ID is available):										
Sponsor amendment reference number*:										
Sponsor amendment date* (enter as DD/MM/YY):										
Briefly summarise in lay language the main changes proposed in this amendment. Explain the purpose of the changes and their significance for the study. If the amendment significantly alters the research design or methodology, or could otherwise affect the scientific value of the study, supporting scientific information should be given (or enclosed separately). Indicate whether or not additional scientific critique has been obtained (note: this field will adapt to the amount of text entered)*:	This amendment is r Change the PI at on Add new participatin	ne recruiting site								
	Specific study									
Project type (select):	Research tissue bank									
					Research database					
Has the study been reviewed by a UKECA-recognised Recommittee (REC) prior to this amendment?:		Yes	5	No						
What type of UKECA-recognised Research Ethics Commi				NHS/HSC REC						
is applicable? (select):					Ministry of I	Defen	ce (MoDRE)			
Is all or part of this amendment being resubmitted to the R Committee (REC) as a modified amendment (i.e. a subst			Yes	6		No				
amendment previously given an unfavourable opinion)? Where is the NHS/HSC Research Ethics Committee (REC) that raviawad	England		Wales	Scotland	No	orthern Irelar			
the study based?:		Yes		No	No		No			
Was the study a clinical trial of an investigational medicinal OR does the amendment make it one?:		Yes	5		No					
EudraCT number*:	2021-003853	-40								
Was this clinical trial of an investigational medicinal processed under the CTIMP combined review service as the Combined Ways of Working (CWoW) pilot)?:			Yes			No				
Did the study receive Pharmacy Assurance?:			Yes			No				
Was the study a clinical investigation or other study of a m does the amendment make it one?:	edical device OR		Yes	5		No				
Did the study involve the administration of radioactive subs requiring ARSAC review, OR does the amendment introdu		Yes	6	No						
Did the study involve the use of research exposures to ion (not involving the administration of radioactive substances amendment introduce this?:		Yes	5	No						
Did the study involve adults lacking capacity OR does the introduce this?:		Yes	3	No						
Did the study involve access to confidential patient informa direct care team without consent OR does the amendmen		Yes	6	No						
Did the study involve prisoners or young offenders who are supervised by the probation service OR does the amender this?:		Yes	3	No						
Did the study involve children OR does the amendment int		Yes	6	No						
Did the study involve NHS/HSC organisations prior to this		Yes	3	No						
Did the study involve non-NHS/HSC organisations OR doe		Yes	3	No						
amendment introduce them?:		England	Т	Wales	Scotland	No	orthern Irelar			
Lead nation for the study:		Yes		No	No		No			
Which nations had participating NHS/HSC organisations p amendment?	rior to this	Yes		Yes	Yes	+	Yes			
Which nations will have participating NHS/HSC organisation		-			-					

Section 2: Summary of change(s)	
What do you want to update?:	Project information
	New site/PI only

Please note: Only use this option when adding a new site or Principal Investigator (PI) to a clinical trial of an investigational medicinal product (CTIMP). You will not be able to make any other changes at the same time.

Please note: Each change being made as part of the amendment must be entered separately. For example, if an amendment to a clinical trial of an investigational medicinal product (CTIMP) involves an update to the Investigator's Brochure (IB), affecting the Reference Safety Information (RSI) and so the information documents to be given to participants, these should be entered into the Amendment Tool as three separate changes. A list of all possible changes is available on the "Glossary of Amendment Options" tab. To add another change, click the "Add another change" box.

	Change 1												
Area of change (select)*:	Researchers												
Specific change (select - only available when area of change is selected first)*:	PI - New PI, or tempo	orary arrangements to cover the absence of a PI											
Further information (free text - note that this field will adapt to the amount of text entered):	Change of PI at Roya Dixit as PI.	Change of PI at Royal Victoria Infirmary, Newcastle: Dr. Michelle Davis will replace Dr. Anand Dixit as PI.											
Applicability:		England	Wales	Scotland	Northern Ireland								
Where are the participating NHS/HSC organisations locate by this change?*:	ed that will be affected	Yes	No	No	No								
Will all participating NHS/HSC organisations be affected by some? (please note that this answer may affect the categ change):	Ą	ll	Some										
			Remove all c	hanges below									

Change 2 Area of change (select)*: Participating Organisations Specific change (select - only available when area of Addition of sites undertaking the same activities as existing sites change is selected first)*: The following sites will join the MAPS-2 trial: 1) South West Acute Hospital and Altnagelvin hospital (Western Health and Social Care Trust), PI: Dr. Breffni Keegan; Further information (free text - note that this field will 2) York Hospital (York and Scarborough Teaching Hospitals NHS Foundation Trust), PI: Dr. adapt to the amount of text entered): Luke Bridge; Antrim Area Hospital (Northern Health and Social Care Trust), PI: Dr. Djamil Vahidassr 3) 4) Broomfield Hospital (Mid & South Essex NHS Hospitals), PI: Dr. Ramanathan Kirthivasan Wales Applicability: England Scotland Northern Ireland Where are the participating NHS/HSC organisations located that will be affected No Yes No Yes by this change?*: Will all participating NHS/HSC organisations be affected by this change, or only All some? (please note that this answer may affect the categorisation for the Some change): Add another change

Declaration by the Sponsor or authorised	delegate
 I confirm that the Sponsor takes responsit I confirm that I have been formally authors 	oility for the completed amendment tool sed by the Sponsor to complete the amendment tool on their behalf
Name [first name and surname]*:	Angela Shone
Email address*:	sponsor@nottingham.ac.uk
Lock for submission	
	vailable when all mandatory (*) fields have been completed. When the button is available, clicking it will ed amendment tool which must be included in the amendment submission. Please ensure that the amendment cathering

After locking the tool, proceed to submit the amendment online. The "Submission Guidance" tab provides further information about the next steps for the amendment.

Section 4: Review bodies for the amendment

		Review bodies																	
	UK wide:				Eng	land a	and Wa	Wales: Scotland:				Northern Ireland:				nd:			
	REC	Competent Authority MHRA - Medicines	Competent Authority MHRA - Devices	ARSAC	Radiation Assurance	UKSW Governance	REC (MCA)	CAG	SddMH	HRA and HCRW Approval	REC (AWIA)	рврр	SPS (RAEC)	National coordinating function	HSC REC	HSC Data Guardians	Prisons	National coordinating function	Category
Change 1:						(Y)				(Y)								Ν	В
Change 2:						(Y)				(Y)								(Y)	New site
Overall reviews for the amend	ment:																		
Full review:						Ν				Ν								Ν	
Notification only:						Υ				Υ								Υ	
Overall amendment type:	No	n-sub	stantia	I, no s	tudy-w	vide re	view r	equire	d										
Overall Category:	В																		