Amendment Tool

v1.6 06 December 2021

Section 1: Project information MAPS-2 Short project title*: IRAS project ID* (or REC reference if no IRAS project ID 290474 is available): MA_15_23 Sponsor amendment reference number*: Sponsor amendment date* (enter as DD/MM/YY): 09 June 2023 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 This amendment consists of the change in PI at one recruiting site, and the addition of three methodology, or could otherwise affect the scientific value new sites (two recruiting sites and one repatriation site). 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)*: Specific study Project type (select): Research tissue bank Research database Has the study been reviewed by a UKECA-recognised Research Ethics Yes No Committee (REC) prior to this amendment?: **NHS/HSC REC** What type of UKECA-recognised Research Ethics Committee (REC) review is applicable? (select): Ministry of Defence (MoDREC) Is all or part of this amendment being resubmitted to the Research Ethics Committee (REC) as a modified amendment (i.e. a substantial amendment Yes No previously given an unfavourable opinion)? England Wales Scotland Northern Ireland Where is the NHS/HSC Research Ethics Committee (REC) that reviewed the study based?: Yes No No No Was the study a clinical trial of an investigational medicinal product (CTIMP) No Yes OR does the amendment make it one?: EudraCT number*: 2021-003853-40 Was this clinical trial of an investigational medicinal product (CTIMP) processed under the CTIMP combined review service (formerly known as Yes No the Combined Ways of Working (CWoW) pilot)?: Did the study receive Pharmacy Assurance?: Yes No Was the study a clinical investigation or other study of a medical device OR Yes No does the amendment make it one?: Did the study involve the administration of radioactive substances, therefore Yes No requiring ARSAC review, OR does the amendment introduce this?:

For office use

QC: No

Did the study involve the use of research exposures to ionising radiation (not involving the administration of radioactive substances) OR does the amendment introduce this?:	Y	′es		No				
Did the study involve adults lacking capacity OR does the amendment introduce this?:	Y	/es		No				
Did the study involve access to confidential patient information outside the direct care team without consent OR does the amendment introduce this?:	Y	és	No					
Did the study involve prisoners or young offenders who are in custody or supervised by the probation service OR does the amendment introduce this?:	Ŷ	′es	No					
Did the study involve children OR does the amendment introduce this?:	Y	′es	Νο					
Did the study involve NHS/HSC organisations prior to this amendment?:	Y	'es	No					
Did the study involve non-NHS/HSC organisations OR does the amendment introduce them?:	Y	/es	No					
	England	Wales	Scotland	Northern Ireland				
Lead nation for the study:	Yes	No	No	No				
Which nations had participating NHS/HSC organisations prior to this amendment?	Yes	Yes	Yes	Yes				
Which nations will have participating NHS/HSC organisations after this amendment?	Yes	Yes	Yes	Yes				

Section 2: Summary of change(s)					
What do you want to undate?	Project information				
hat do you want to update?:	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 tempor	rary arrangements	to cover the abse	nce of a PI		
Further information (free text - note that this field will adapt to the amount of text entered):	The Principal Investiga from Dr. Anna Bahk to	•		y Hospitals NHS T	rust has changed	
Applicability:		England	Wales	Scotland	Northern Ireland	
Where are the participating NHS/HSC organisations locate by this change?*:	d 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 categor change):	A	AII	So	Some		
				Remove all c	hanges below	

	Change 2									
Area of change (select)*:	Participating Organisations									
Specific change (select - only available when area of change is selected first)*:	Addition of sites undertaking the same activities as existing sites									
Further information (free text - note that this field will adapt to the amount of text entered):	Three new sites are joining the trial: 1) Ulster Hospital (South Eastern Health and Social Care Trust), PI: Dr. Mark Bowman. 2) Royal Shrewsbury Hospital / Princess Royal Hospital (Shrewsbury and Telford Hospital NHS Trust), PI: Dr. Indranil Mukherjee. 3) Trafford General Hospital (Manchester University NHS Foundation Trust), PI: Mr Terence Kelly.									
Applicability:	England	Wales	Scotland Northern Irela							
Where are the participating NHS/HSC organisations located by this change?*:	Yes	No	No Yes							
Will all participating NHS/HSC organisations be affected by this change, or only some? (please note that this answer may affect the categorisation for the change):		A		Some						
				Add anot	her change					

Section 3: Declaration(s) and lock for submission

Declaration by the Sponsor or authorised delegate

• I confirm that the Sponsor takes responsibility for the completed amendment tool

• I confirm that I have been formally authorised 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

Please note: This button will only become available when all mandatory (*) fields have been completed. When the button is available, clicking it will generate a locked PDF copy of the completed amendment tool which must be included in the amendment submission. Please ensure that the amendment tool is completed correctly before locking it for submission.

Lock for submission

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	England and Wales:			Scotland:				Northern Ireland:				
	REC	Competent Authority MHRA - Medicines	Competent Authority MHRA - Devices	ARSAC	Radiation Assurance	UKSW Governance	REC (MCA)	CAG	SHIMPPS	HRA and HCRW Approval	REC (AWIA)	BPP	PS (RAEC)	Vational coordinating function	SC REC	HSC Data Guardians	risons	National coordinating function	Categor
Change 1:		$\odot \ge$	$\odot \ge$	A		⊃ (Y)		0	I	エ (Y)		<u> </u>	0	Z	I	工	<u> </u>	Z N	B
Change 2:						(Y)				(Y)								(Y)	New sit
Overall reviews for the amendr	ment:																		
Full review:						Ν				Ν								N	
Notification only:						Y				Υ								Y	
Overall amendment type:	Nor	n-subst	tantial,	no st	udy-wi	de rev	view re	quirec											
Overall Category:	В																		