v1.6 06 December 2021

Short project title*:	The GenOMICC Stud	dy									
IRAS project ID* (or REC reference if no IRAS project ID is available):	189676 (Scotland) 2	69326 (Rest of UK)									
Sponsor amendment reference number*:	SA_014 (England) a	nd SA_011 (Scotland)									
Sponsor amendment date* (enter as DD/MM/YY):	29 October 2024										
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)*:	We propose to expand our objectives, by widening our inclusion criteria and introducing a s study, ACUTE. ACUTE follows the exact methodolgy of GenOMICC but requires a small an of additional blood. We have also modernised the font within the protocol and made minor e both within the protocol and across the study documents, including clarifying the withdrawal process and defining the role of the consultee in the telephone script. The changes are defit the summary of changes attachment.										
				Specific stud	dy						
Project type (select):				Research tiss	sue bank						
				Research dat	abase						
Has the study been reviewed by a UKECA-recognised Resea Committee (REC) prior to this amendment?:	arch Ethics	Ye	es		No						
· //	NHS/HSC REC										
What type of UKECA-recognised Research Ethics Committee applicable? (select):	e (REC) review is			Ministry of De	efence (MoDRE)						
Is all or part of this amendment being resubmitted to the Resc Committee (REC) as a modified amendment (i.e. a substan previously given an unfavourable opinion)?	Ye	Yes No									
Where is the NHS/HSC Research Ethics Committee (REC) tl	hat reviewed the	England	Wales	Scotland	Northern Irela						
study based?:	nat reviewed the	No	No	Yes	No						
Was the study a clinical trial of an investigational medicinal p OR does the amendment make it one?:	product (CTIMP)	Ye	es	No No							
Was the study a clinical investigation or other study of a med does the amendment make it one?:	lical device OR	Ye	es								
Did the study involve the administration of radioactive substa requiring ARSAC review, OR does the amendment introduce		Ye	es		No						
Did the study involve the use of research exposures to ionisin involving the administration of radioactive substances) OR do amendment introduce this?:		Ye	es	ı	No						
Did the study involve adults lacking capacity OR does the an introduce this?:	nendment	Yes No									
Did the study involve access to confidential patient informatic direct care team without consent OR does the amendment in		Ye	es	No							
Did the study involve prisoners or young offenders who are in supervised by the probation service OR does the amendmen		Ye	Yes No								
Did the study involve children OR does the amendment introd	duce this?:	Ye	es		No						
Did the study involve NHS/HSC organisations prior to this am	nendment?:	Ye	es		No						
Did the study involve non-NHS/HSC organisations OR does to introduce them?:	the amendment	Ye	es	ı	No						
		England	Wales	Scotland	Northern Irela						
Lead nation for the study:		No	No	Yes	No						
Which nations had participating NHS/HSC organisations prio amendment?	or to this	Yes	Yes	Yes	Yes						
Which nations will have participating NHS/HSC organisations	s after this	Yes	Yes	Yes	Yes						

Section 2: Summary of change(s)

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)*:	Study Design

Specific change (select - only available when area of change is selected first)*:	Inclusion/exclusion criteria - Significant change (e.g. to age range) likely to affect safety or scientific value of study							
Further information (free text - note that this field will adapt to the amount of text entered):	Changes to the inclusion criteria to include all infections and addition of pre-term babies							
Applicability:	England	Wales	Scotland	Northern Ireland				
Where are the participating NHS/HSC organisations located this change?*:	Yes	Yes	Yes 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):		All Some						
	Remove all o	changes below						

	Change 2								
a of change (select)*: Study Documents									
Specific change (select - only available when area of change is selected first)*:	Other significant change to study documents (e.g. information sheets, consent forms, questionnaires, letters) that can be implemented within existing resource in place at participati organisations - Please specify in the free text below								
Further information In particular, please describe why this change can be implemented within the existing resource in place at the participating organisations (free text - note that this field will adapt to the amount of text entered)*									
Applicability:		England	Wales	Scotland	Northern Ireland				
Where are the participating NHS/HSC organisations located t this change?*:	hat will be affected by	Yes	Yes	Yes	Yes				
Will all participating NHS/HSC organisations be affected by the some? (please note that this answer may affect the categorise	0,	All Some							
				Remove all	changes below				

Change 3									
Area of change (select)*:									
Specific change (select - only available when area of change is selected first)*:	Other minor change to study documents (e.g. information sheets, consent forms, questionnal letters) that can be implemented within existing resource in place at participating organisation Please specify in the free text below								
Further information In particular, please describe why this change can be implemented within the existing resource in place at the participating organisations (free text - note that this field will adapt to the amount of text entered)*									
Applicability:		England	Wales	Scotland	Northern Ireland				
Where are the participating NHS/HSC organisations located this change?*:	Yes	Yes	Yes	Yes					
Will all participating NHS/HSC organisations be affected by the some? (please note that this answer may affect the categorise	All Some								
		Remove all o	changes below						

Change 4										
Area of change (select)*:	Study Design									
	Other significant change to study design that can be implemented within existing resource in place at participating organisations - Please specify in the free text below									
Further information In particular, please describe why this change can be implemented within the existing resource in place at the participating organisations (free text - note that this field will adapt to the amount of text entered)*										
Applicability:		England	Wales	Scotland	Northern Ireland					
Where are the participating NHS/HSC organisations located that this change?*:	at will be affected by	Yes	Yes	Yes	Yes					
Will all participating NHS/HSC organisations be affected by this some? (please note that this answer may affect the categorisa	All 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]*:	Jo-Anne Robertson
Email address*:	genomicc@roslin.ed.ac.uk andjo-anne.robertson@ed.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

Please note: This section is for information only. Details in this section will complete automatically based on the options selected in Sections 1 and 2.

		Review bodies																	
	UK wide:				Eng	land a	nd Wa	les:		Scot	land:		Northern Ireland:						
	REC	Competent Authority MHRA - Medicines	Competent Authority MHRA - Devices	ARSAC	Radiation Assurance	UKSW Governance	REC (MCA)	CAG	HMPPS	HRA and HCRW Approval	REC (AWIA)	РВРР	SPS (RAEC)	National coordinating function	HSC REC	HSC Data Guardians	Prisons	National coordinating function	Category:
Change 1:	Υ					(Y)				(Y)				(Y)				(Y)	Α
Change 2:	Υ					Υ				Υ				Υ				Υ	С
Change 3:	N					(Y)				(Y)				(Y)				(Y)	С
Change 4:	Υ					Υ				Υ				Υ				Υ	С
Overall reviews for the amendmen	t:	•																•	
Full review:	Υ					Υ				Υ				Υ				Υ	
Notification only:	N					N				N				N				N	
Overall amendment type:	Su	bstanti	al																
Overall Category:	Α																		