Modification Tool

v2.1 24 April 2026

For office use

QC: No

Section 1: Project information

Short project title*:	SWEET
IRAS project ID* (or REC reference if no IRAS project ID is available):	330129
Sponsor modification reference number*:	NSM24
Sponsor modification date* (enter as DD/MM/YY):	07 October 2026
Briefly summarise in lay language the main changes proposed in this modification. Explain the purpose of the changes and their significance for the project. If the modification 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)*:	Update to the following document SWEET Follow Up Cover Letter From SWEET_FollowUpCoverLetter_v2.0_17Mar2025 To sweet_followupcoverletter_v3.0_17Sept2026 Changes include * Removal of 6 and 12 month timepoint * Highlighting to particpants this is the final questionnaire * Administrative/ wording changes tohighlight the importance of the final questionaire

Project type (select):	**Specific project** Research tissue bank Research database			
Has the project been reviewed by a UKECA-recognised Research Ethics Committee (REC) prior to this modification?:	**Yes**	No		
What type of UKECA-recognised Research Ethics Committee (REC) review is applicable? (select):	**NHS/HSC REC** Ministry of Defence (MoDREC)			
Where is the NHS/HSC Research Ethics Committee (REC) that reviewed the project based?:	England	Wales	Scotland	Northern Ireland
	Yes	No	No	No
Was the project a clinical trial of an investigational medicinal product (CTIMP) OR does the modification make it one?:	Yes		**No**	
Was the project a clinical investigation or other study of a medical device OR does the modification make it one?:	Yes		**No**	
Was the project a performance evaluation study of an In Vitro Diagnostic (IVD) medical device^ (including as a companion diagnostic in a clinical trial of an investigational medicinal product)?:	Yes		**No**	
Does the modification make the project a performance evaluation study of an In Vitro Diagnostic (IVD) medical device^ (including as a companion diagnostic in a clinical trial of an investigational medicinal product)?:	Yes		**No**	
^ IVD medical devices are tests used on biological samples, such as tissues, blood or urine, to determine the status of a person's health. This may be a reagent, reagent product, calibrator, control material, kit, instrument, apparatus, equipment or system, whether used alone or in combination				
Did the project involve the administration of radioactive substances, therefore requiring ARSAC review, OR does the modification introduce this?:	Yes		**No**	
Did the project involve the use of research exposures to ionising radiation (not involving the administration of radioactive substances) OR does the modification introduce this?:	Yes		**No**	
Did the project involve adults lacking capacity OR does the modification introduce this?:	Yes		**No**	
Does the project involve access to confidential patient information outside the direct care team without consent OR does the modification introduce this?: (e.g. the project relies upon section 251 support in England and Wales, or equivalent in Scotland to set aside the common law duty of confidentiality)	Yes		**No**	
Does the project involve prisoners or young offenders who are in custody or supervised by the probation service OR does the modification introduce this?:	Yes		**No**	
Does the project involve children OR does the modification introduce this?:	Yes		**No**	
Did the study involve NHS/HSC organisations prior to this modification?:	**Yes**		No	
Did the study involve non-NHS/HSC organisations OR does the modification introduce them?:	Yes		**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 modification?	**Yes**	**Yes**	**Yes**	**Yes**
Which nations will have participating NHS/HSC organisations after this modification?	**Yes**	**Yes**	**Yes**	**Yes**

Section 2: Summary of change(s)

Please note: Each change being made as part of the modification must be entered separately. For example, if a modification 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 modification tool as three separate changes. A list of all possible changes is available on the "Glossary of Options" tab. To add another change, click the "Add another change" box.

Change 1	
Area of change (select)*:	Project documents
Specific area of change (select - only available when area of change is selected first)*:	Other non-significant change to project documents that will have additional resource implications for participating organisations - please specify in the free text below
Further information: explain what the change is, why the change is being made and any possible impact on participants/carers. In particular, please describe the additional resource arrangements that participating organisations will need to have in place to implement this change (free text - note that this field will adapt to the amount of text entered)*:	Update to the following document SWEET Follow Up Cover Letter From SWEET_FollowUpCoverLetter_v2.0_17Mar2025 To sweet_followupcoverletter_v3.0_17Sept2026 Changes include * Removal of 6 and 12 month timepoint * Highlighting to particpants this is the final questionnaire * Administrative/ wording changes tohighlight the importance of the final questionaire

Applicability:	England	Wales	Scotland	Northern Ireland
Where are the participating NHS/HSC organisations located that will be affected by 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	

Add another 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 modification tool
- I confirm that I have been formally authorised by the Sponsor to complete the modification tool on their behalf

Name [first name and surname]:*	Christopher Price
Email address:*	tnu-tr.sponsormanagement@nhs.net

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 modification tool which must be included in the modification submission. Please ensure that the modification tool is completed correctly before locking it for submission.

Lock for submission

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

Section 4: Review bodies for the modification

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:						England and Wales:				Scotland:				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)	PBPP	SPS (RAEC)	National coordinating function	HSC REC	HSC Data Guardians	Prisons	National coordinating function	Category:
Change 1:						(Y)				(Y)				(Y)				(Y)	A
Overall reviews for the modification:																			

Full review:						N				N				N				N
Notification only:						Y				Y				Y				Y
Overall modification type:	Minor Modification																	
UKSW Governance:	NHS/HSC review is not required																	
Overall Category:	A																	

Please note: The modification tool has been revised to provide updated classifications in line with the new regulations. However, IRAS has not been updated and will still require you to submit using the old classifications. This section provides guidance on which option to select in order to ensure the modification is processed correctly:

Your modification should be submitted as a **non-substantial, no study-wide review required amendment**