## **Amendment Tool**

v1 2 11 Jun 2020

For office use QC: No

Section 1: Project information											
Short project title*:	OPD										
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):	10 August 2020										
Summary of amendment including justification*:	Addition of site (non-ctimp study)										
		•	Specific study								
Project type:		0	Research tissu	ue bank							
		0	Research data	abase							
Has the study been reviewed by a UKECA-recognised Reso Committee (REC) prior to this amendment?:	•	Yes	C	No No							
	•	NHS/HSC RE	C								
What type of UKECA-recognised Research Ethics Committ is applicable?:	O Ministry of Defence (MoDREC)										
Is all or part of this amendment being resubmitted to the Re Committee (REC) as a modified amendment?	0	Yes	•	No No							
Where is the NHS/HSC Research Ethics Committee (REC)	that reviewed	England	Wales	Scotland	Northern Ireland						
the study based?:  Was the study a clinical trial of an investigational medicinal	product (CTIMP)	•	0	0	0						
OR does the amendment make it one?:		0	Yes	•	No No						
Was the study a clinical investigation or other study of a me does the amendment make it one?:	edical device OR	0	Yes	•	No No						
Did the study involve the administration of radioactive substrequiring ARSAC review, OR does the amendment introduce		0	Yes	•	No No						
Did the study involve the use of research exposures to ionis involving the administration of radioactive substances) OR amendment introduce this?:		0	Yes	•	) No						
Did the study involve adults lacking capacity OR does the a introduce this?:	mendment	O Yes ® No									
Did the study involve access to confidential patient informat consent OR does the amendment introduce this?:	tion without	0	) No								
Did the study involve prisoners OR does the amendment in	troduce this?:	O Yes ® No									
Did the study involve NHS/HSC organisations prior to this a	Did the study involve NHS/HSC organisations prior to this amendment?:										
Did the study involve non-NHS/HSC organisations OR does introduce them?:	0	Yes	•	No No							
		England	Wales	Scotland	Northern Ireland						
Lead nation for the study:		•	0	0	0						
Which nations had participating NHS/HSC organisations pramendment?	ior to this	Ø									
Which nations will have participating NHS/HSC organisation amendment?	ns after this	Ø									

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 Changes" tab. To add another change, tick the "Add another change" box.

	Change 1												
Area of change (select)*:													
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):	dition of University l	Hospital Coventry	and Warwickshire	PI Lara Beatriz									
Applicability:	England	Wales	Scotland	Northern Ireland									
Where are the participating NHS/HSC organisations located that this change?*:	v												
Will all participating NHS/HSC organisations be affected by this c some?:	•	All	(	Some									
				Add another cha	nge:								

Section 3: Declaration(s) and lock for submission
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## 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]*:	Dalbir Kaur
Email address*:	d.kaur@bham.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 PDF copy of the completed amendment tool that can 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, refer to the "Submission Guidance" tab for further information about the next steps for the amendment.

## Section 4: Review bodies for the amendment

Please note: This section is for inform	matio	n only	. Deta	ils in th	nis sec	tion w	ill com	plete a	automa	atically	based	d on th	ne opti	ons se	lected	in Se	ctions	1 and	2.
	Review bodies																		
	UK wide:					Eng	land a	nd Wa	ales:	Scotland:				Northern Ireland:					
	REC	Competent Authority JHRA - Medicines	Competent Authority MHRA - Devices	ARSAC	adiation Assurance	UKSW Governance	REC (MCA)	AG	HMPPS	HRA and HCRW Approval	REC (AWIA)	ВРР	SPS (RAEC)	National coordinating function	HSC REC	HSC Data Guardians	risons	National coordinating function	Category:
Change 1:	K	0 2	02	ď	ď	(Y)	Ľ	Ö		(Y)	<u>K</u>		S				۵		New site

Overall reviews for the amendmen	nt:															
Full review:						N				N						
Notification only:						Υ				Υ						
Overall amendment type:	Non-substantial, no study-wide review required															
Overall Category:	Ne	w site														