FORM FRM4272/4.1 - Continuous Improvement Event A3

1. Improvement Details

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EFFECTIVE: 20/09/17



Title:	Research Application process
Date:	23-24 November 2017
Venue:	Cambridge
Sponsor:	ODT
Team Leader:	Janine Stockdale
Facilitator / Sensei:	Janine Stockdale

Scoping Complete? Yes

Initial Risk Assessment

Please see Risk Assessment in Controlled Documents, under CC/8533

Outcome of Assessment:

Needs Improvement

Change Control (CC) Required?

Yes. Number CC/8533

Team members:

Gavin Pettigrew, Vicky Gauden, Cat MacDonald, Liz Armstrong, Maggie Stevens, Janine Stockdale, Maria McGee, Nick Watson, Yomi Adegbaju, Mick Stokes, Kourosh Saeb-Parsy, Jenny Mehew, Alia Rashid, Bill Scott

Post Activity Reviews

First Sustainment Review Date:

dd/mm/yy

Second Sustainment Review Date:

dd/mm/yy

Final Sustainment Review & Closure Date:

dd/mm/yy

Compl	etion	Deta	ails:
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S	ponso	or	Confirms	Activity	Cycle	Complete?
_	TO MAKE	and the state of t				

On closure:

Sign off: Closure Date: _ _ / _ _ /

Have you achieved what you set out to do? Y/N

Yes – when will you return to review this process?

No – what are you planning to do as part of PDCA? Eg another Improvement A3

2. Reason for Action

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Context:

The ODT research application process has been extensively reviewed and changed over the last 2 years. Changes to internal and external application processes resulted in some unforeseen delays.

Following internal and external stakeholder feedback, the application process is now to be subject to lean review.

Link your performance issues to the relevant improvement themes below with a short general statement. Where themes are not applicable consider potential negative impact.

A) <u>Impact on people</u>

Delays for researchers starting projects due to lengthy application processes.

Impact on donor families who are asked for consent for the research.

Internal stakeholders are impacted by an inconsistent approach being sought for advice and guidance.

In scope:

Entire application process

B) Quality of products or services

Application process needs to be streamlined and consistent to ensure the quality of the application process service.

Out of scope:

External approvals process:

HRA

HTA

REC

Local Trust MoU for researchers

C) Cost or productivity

Productivity is affected by the complexity and length of time that each research project takes to facilitate.

Costs can be occurred by the researchers due to delays in starting. Funding may be revoked.

D) <u>Delivery or timeliness</u>

Delays starting projects due to lengthy application processes resulting reputational issues for NHSBT.

3. Current and Target Performance

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CURRENT

A) Impact on people

Delays for researchers starting projects due to lengthy application processes.

Impact on donor families who are asked for consent for the research.

Internal stakeholders are impacted by an inconsistent approach being sought for advice and guidance

B) Quality of product or services

Application process needs to be streamlined and consistent to ensure the quality of the application process service.

C) Cost or productivity

Productivity is affected by the complexity and length of time that each research project takes to facilitate.

Costs can be occurred by the researchers due to delays in starting. Funding may be revoked.

D) <u>Delivery or timeliness</u>

Delays starting projects due to lengthy application processes resulting reputational issues for NHSBT.

TARGET

A) Impact on people

No delays

Positive experience for researchers and internal stakeholders

B) Quality of product or services

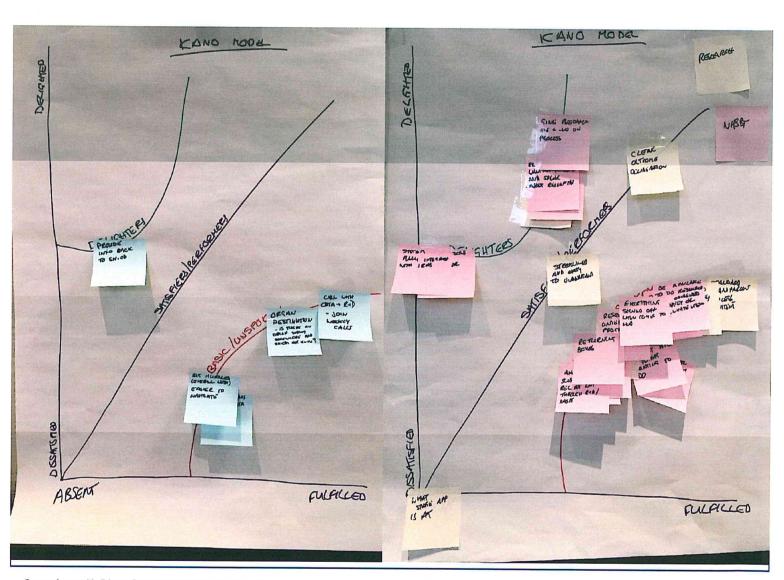
Consistent quality throughout the research application process.

C) Cost or productivity

Productivity increases due to streamlined processes.

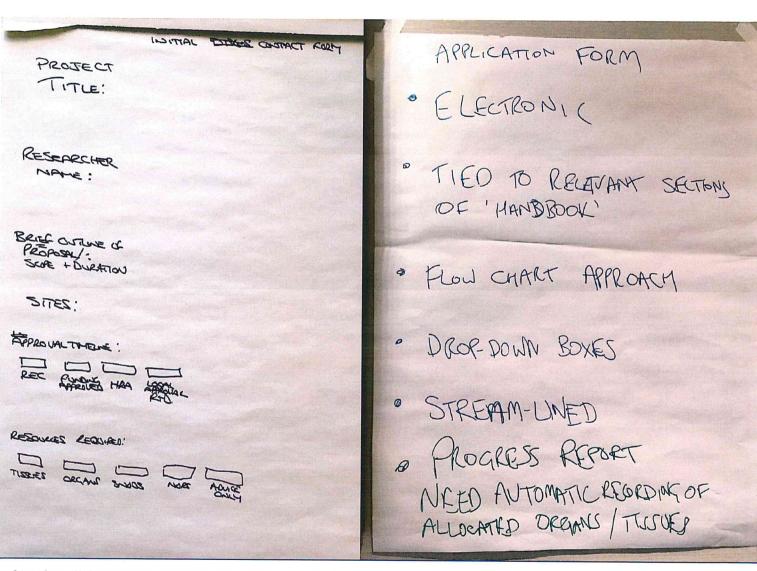
D) <u>Delivery or timeliness</u>

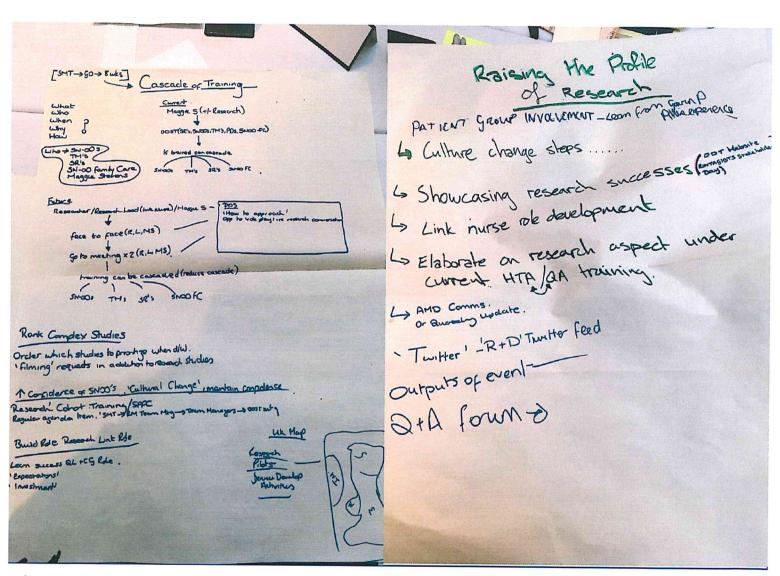
Delivery is improved and timely.



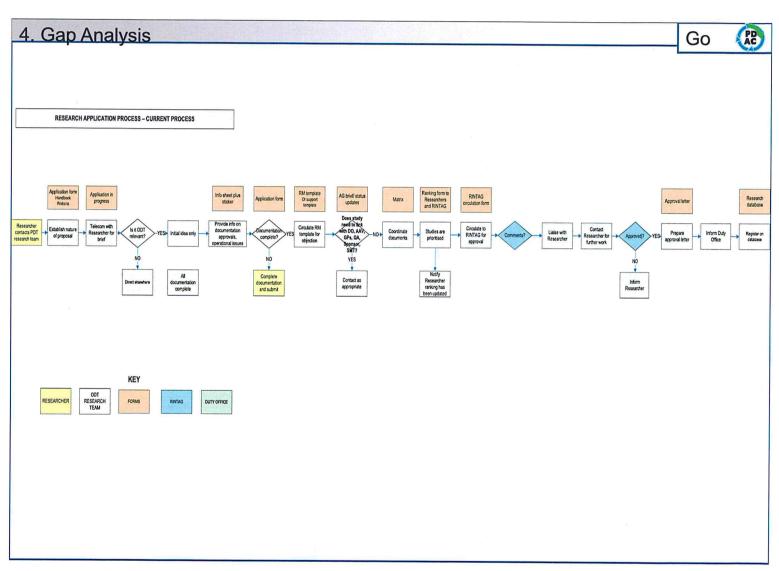
Cross referenced in Primary Document: SOP3921 Creation and Management of Continuous Improvement Documents

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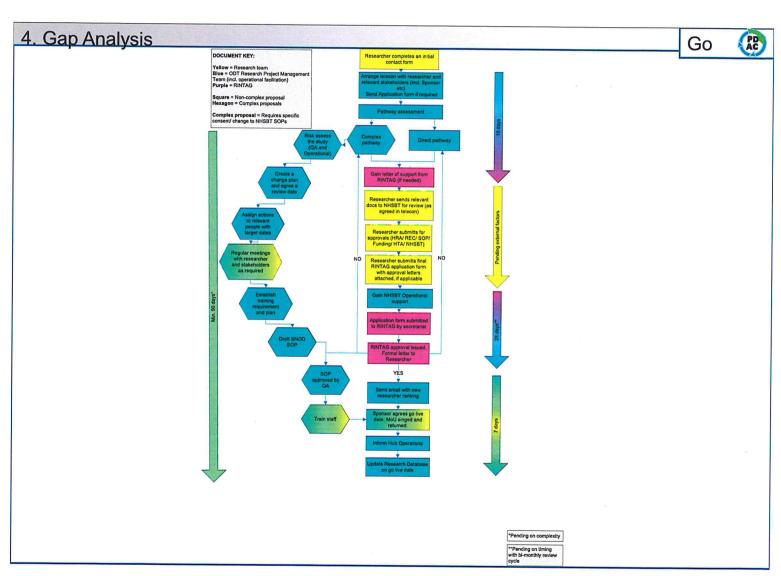




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RINTAG(18)1

Solution	Action	Who	By When	R/G
	New approach presented to and approved by RINTAG	MM	May 2018	
	Development of ODT Research KPIs	MM	May 2018	
	Develop and trial electronic forms	MM/ All	March 2018	
	Discussion w PDS team reg. roll out of training	MS	March 2018	G
	Website, showcase success	MM/ LA	May 2018	
	Communicate purpose of RINTAG and ODT Research	MM/ MS	May 2018	
	Create FAQs for RECs	BS/ K-SP	May 2018	
	Cost recovery business case	LA	May 2018	
	Communicate the new process, including instructions and a link to the Cognito forms, to external stakeholders (researchers and transplant surgeons, Sponsors)	ММ	May 2018	
	Control the new process on QPulse and update the website with the new documents	MM	May 2018	
	Create an animation video for research community and put this on the website and circulate to RMs and SNODs for information	MM/ MS	March 2018	G
	Training on new controlled docs as appropriate (MPD1029)	MM	May 2018	
	Confirm Information Governance risk regarding the use of Cognito forms	MM/ LA	May 2018	

