

Module code

Supervisor's name

Second Marker's name

participants:

No participants will be required for this research.

Start Date of project

Short description of project, including research methods and selection of any

Non-experimental research will be conducted with quantitative data analysis methods.

Dr Gregory Epiphaniou 01 August 2016

LD0718

Dr Haider Al-Khateeb

Ethical considerations in the research project	YES	NO	
1. Does your research involve an external organisation or partner e.g. NHS, School etc.			
2. Does your research involve human participants?		\boxtimes	
3. If yes to Q.2, will you inform the participants about the research?			
4. Will you obtain their consent using the standard consent form?			
5. Is any deception involved?			
6. Do any participants constitute a 'vulnerable group' (see definition of Vulnerable People)			
7. Will the research involve the following information?			
Commercially sensitive		\boxtimes	
Personally sensitive		\boxtimes	
Politically sensitive	Ш	\boxtimes	
8. Is the research likely to cause any significant environmental impacts?		\boxtimes	
9. Are there likely to be any risks for you or for the participants in your research?		\boxtimes	
10. If yes [to 5, 6, 7, 8 or 9 above] have you identified steps to address the issues?			
Statement by researcher			
This statement should explain how any issues identified in the answers to the above questions will be addressed and what steps will be taken to mitigate such risks or adverse impact			
There are no issues that needs to be addressed.			

I have read the University and the Faculty Ethics Policy and Procedures and confirm that the answers I have given above are correct. Where issues arise under items 5, 6, 7. 8 or 9 [above] I have described in writing how I intend to approach these issues in the research. Raidey exam



Section Two: Approval		
[The form is reviewed by the supervisor and second marker. Approval maybe given by either for green projects; amber projects must be approved by the second marker. Red projects must be referred to the Faculty Research Ethics Committee.]		
Red : Vulnerable participants, sensitive data, risks to participants or researchers, NHS, Amber: Human participants, environmental issues, commercially sensitive information Green : No participants involved, no sensitive data, etc. For full definitions see section on Risk Categories in the Engineering and Environmental Procedures.	, etc.	
Ethical approval		
Green - Ethical approval is given without conditions	\boxtimes	
Amber - Ethical approval is given with the following conditions		
Information to be provided to all participants		
Participant consent to be obtained using the standard Research Participant		
Consent Form or otherwise in accordance with Faculty procedures		
Data to be stored and destroyed securely in accordance with University		
guidelines • Adherence to Data Protection Act (DPA)		
 Adherence to Data Protection Act (DPA) Anonymity to be provided to participants 		
Commercial confidentiality to be provided to organisations(s)		
Other (please state):		
Red - Project is referred to FREC for approval		
Name of Approver		
Signature		
Date 01 August 2016		
Outcome of FREC referral – Decision, minute and date of meeting, or signature two signatories, one of whom is a member of FREC.	res of	