

## **Clinician Perceptions Participation Information Sheet**

The 2nd Sprint National Anaesthesia Project (SNAP-2): Epidemiology of Critical Care provision after Surgery (EpiCCS)

The Sprint National Anaesthesia Projects (SNAPs) are 'snapshot' evaluation studies of clinical activity and patient-centred outcomes that are important and relevant to both patients and anaesthetists. SNAP-1 was a two-day evaluation of patient reported outcomes after anaesthesia conducted in May 2014 and over 95% of NHS hospitals in all 4 devolved UK nations participated in that study.

We are now conducting the second SNAP research study (SNAP-2: EpiCCS), a prospective observational cohort study looking at postoperative Critical Care in hospitals throughout the UK, and would like to invite you to take part. Please take time to read the following information carefully. If there is anything that is not clear or if you would like more information, please contact the Local Lead Investigators (details can be found at the bottom of this information sheet).

#### What is the purpose of the study?

Postoperative complications, such as post-operative nausea and vomiting, infections and death, can vary in their incidence and severity. It has been suggested that one way to reduce complications is to admit patients to Critical Care postoperatively, for more intensive nursing support, or advanced interventions, etc. However, despite national guidelines and recommendations, we know that that there is variation in which patients are cared for in Critical Care in different institutions and countries. We are conducting this study to try and uncover some of the reasons for these findings. We also hope to find out whether Critical Care after surgery shows a beneficial effect on morbidity and mortality after surgery. This is particularly important as the thresholds for critical care admission are not clear from the evidence, and opinions are not consistent between clinicians or hospitals.

#### Why have I been invited?

Part of our study is looking at reasons why some high-risk patients might not be admitted to a Critical Care bed immediately following surgery. We are therefore interested in seeing how clinicians (anaesthetists, surgeons and intensivists) decide where patients should go for postoperative care.

#### Do I have to take part?

Completing the questionnaire is entirely voluntary and it is up to you to decide whether you do so. You are free to withdraw at any time. If you choose not to take part or to withdraw your participation, we ask that you provide a reason why, so that we can better account for participant withdrawal during data analysis.

#### What will I have to do?

We will ask you to complete the short questionnaire on page 3. You will only have to complete this questionnaire once during the SNAP-2 study week. By completing this questionnaire, you are giving permission for us to keep the information you provide and analyse it for our study.



#### What are the possible disadvantages and risks of taking part?

There are no major disadvantages or risks of taking part. Filling in the questionnaires should be straightforward and does not take up much time. We would like to reassure you that your responses will be used in confidence, without risk of litigation or reprisal. All data will be anonymised so there is no risk of you being identified.

What are the possible benefits of taking part?

We cannot promise the study will help you directly in caring for your patients but we hope that the information you provide will help improve the quality of clinical care delivered in the future. You will be able to get feedback on the findings of the study via the SNAP-2 website (see details below).

Will my taking part in the study be kept confidential?

Yes. The study will abide by strict information governance and confidentiality procedures. The information you provide will be anonymised for analysis.

What will happen to the answers provided in the questionnaires?

The Local Lead Investigator will transfer your responses from the paper questionnaires onto a computer database. Data analysis will be performed by researchers based at the Health Services Research Centre and University College Hospital in London. Your information will be kept securely for 10 years in order for long-term outcomes to be accurately studied.

What will happen to the results of the study?

The results from the study will be published in scientific manuscripts and on the SNAP-2 website. No references will be made that could link you personally to the study.

Who is organising and funding the study?

The study is being funded by the National Institute of Academic Anaesthesia (NIAA), the Association of Anaesthetists of Great Britain & Ireland, the NIAA Health Services Research Centre (HSRC) & the Royal College of Anaesthetists. It is being led by the NIAA HSRC.

Who has reviewed the study?

This study has been reviewed and approved by the South Central - Berkshire B Research Ethics Committee (REC reference: 16/SC/0349) on behalf of the Health Research Authority.

Further information and contact details

Website: http://www.niaa.org.uk/SNAPs

Study email address: snap2@rcoa.ac.uk

Local lead name and contact details:

Appendix 4 – Clinician Perception Substudy CRF, Version 1.0. Last amended 01/02/2017

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# Clinician Perceptions to risk stratification and postoperative care

We are surveying anaesthetists, surgeons and intensivists to explore their approaches to risk stratification and decision-making surrounding postoperative care.

This should only be completed once by each clinician during the week of the study.

About You							
1.1. What is your specialty? (tick all that apply)		Anaesthetist	Surgeon	□Intensivist		<b>Commented [A1]:</b> We are also encouraging Interventional Radiologists, Endoscopists and Cardiologists to participate,	
1.1a. If you are a surgeon o select more than one)?	or anaesthetist, which	ch of the following best	t describes your p	primary workload (you may		please ask them to select "Surgeon".	
□Cardiothoracic surgery	□Ger	□General surgery (including GI surgery and breast surgery)				describes your primary workload, please write down a free text response. We will include the option for "Other" and space for free-text on the online webtool.	
□ Neurosurgery	□Ora	$\Box$ Oral and maxillofacial surgery					
□ Otolaryngology	□Plas						
□Trauma and Orthopaedic	cs 🗌 Uro	logy					
□Vascular surgery							
1.2. What is your grade?	Consultant	□St	aff & Associate s	pecialist			
	ST3-7 trainee or Tru	ist grade equivalent					
□ Core/Foundation trainee or Trust grade equivalent							
1.3. Date of completion of questionnaire://							
1.4. How many years have you been	n qualified as a doc	tor?					
About your views on risk pred	diction and Criti	cal Care					
2.1. What predicted risk of 30-day mortality would you set as a minimum threshold for admitting patients to critical care immediately after surgery? (i.e. if the predicted risk was above this level you would seek a planned critical care admission)						<b>Commented [A3]:</b> We understand that this question may seem very subjective. That is the intention. We want to find out what the clinicians' "gut" feelings are about how risky a case might need to be before they would consider sending a	
□<1% □1-2.5%	□2.6-5%	□5.1-10%	□10.1-50%	□>50%		patient to Critical Care.	
2.2. What risk of 30-day mortality v	vould you consider	to be "high-risk" for a p	patient undergoi	ng surgery?			
Please indicate a value between 0 a	and 100%:						
2.3. In general, do you feel that there is enough critical care bed capacity at your institution for postoperative patients that need it?							
□Y □N	□Not sure						

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Please elaborate if you would like to contribute a free text response to this question:

2.4. Do you think the they were admitted	at high-risk pati onto a general		
ΠY	□N	□Not sure	
Please elabo	orate if you wou	Id like to contribute a free text response to this question:	
2.5. Would you proc surgery?	ceed with surge	ry for a high-risk patient if you knew there was no critical care capacity on the day of	
ΠY	□N		
Please elabo	orate if you wou	Ild like to contribute a free text response to this question:	
2.6. Do pathways ex postoperatively?	kist within your	institution for certain operations mandating that patients be admitted to critical care	<b>Commented [A4]:</b> An example of this might be: all cardiopulmonary bypass patients are admitted to critical care postoperatively as a matter of routine.
ΠY	□N	□Not sure	
2.7. Are there specif vs. emergency surge	fic types of surg ery, open laparc		
2.8. What do you fe important) to <b>6</b> (lea:	el is the benefit st important), if		
ncreased monitor	oring	Higher nurse:patient ratio	
bility to deliver	complex thera	pies Higher doctor:patient ratio	
Consultant Inten	isivist input		

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### **Risk stratification**

# How frequently do you use the following risk stratification tools in your perioperative practice?

(Please circle **0** to **4**, where: **0** = Never, **1** = Occasionally, **2** = Usually and **3** = Always)

1. American Society of Anaesthesiologists Physical Status (ASA-PS)	Never 0	1	2	3 always
2. Charlson Comorbidity Index	Never 0	1	2	3 always
3. Revised Cardiac Risk Index (RCRI)	Never 0	1	2	3 always
4. Surgical Risk Scale (SRS)	Never 0	1	2	3 always
5. POSSUM and/or associated derivatives <sup>1</sup>	Never 0	1	2	3 always
6. Surgical Outcome Risk Tool (SORT)	Never 0	1	2	3 always
7. Cardiopulmonary exercise testing (CPET)	Never 0	1	2	3 always
8. EuroSCORE	Never 0	1	2	3 always
9. Duke Activity Status Index or other Metabolic Equivalents (MET) Scoring sy	ystem Never 0	1	2	3 always
10. New York Heart Association Functional Classification	Never 0	1	2	3 always
11. Acute Physiology & Chronic Health Evaluation II (APACHE-II)			2	3 always
12. Sequential Organ Failure Assessment (SOFA)		1	2	3 always

<sup>1</sup> Physiological and Operative Severity Score for the enUmeration of Mortality and Morbidity, including p-POSSUM, v-POSSUM, etc.

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