



Postoperative vasopressor usage: a prospective international observational

study

CRF1		PATIENT IDENTIFICATION NUMBER:	
PRE-OPERATIVE		DATE OF SURGERY:	
Month and year of birth			
Sex [m/f]			
Height [cm]			
Veight [kg]			
	kwood): point 0 to 9. (Will be explained in	final CRF)	
Previous medical history:			
Coronary Artery Dis Cerebrovascular Dis			
Peripheral vascular			
Atrial fibrillation: Y/	N		
Heart failure: Y/N Hypertension: Y trea	ated and controlled, Y treated but not controlled, No		
	ulin/managed without insulin/None		
Chronic liver disease			
	disease: COPD/other/None ppression: HIV/other/none		
	ase: No/Yes/Yes and receives renal replacement ther	ару	
Long-term steroid u			
Recent/current trea Regular medications	Itment for cancer (including chemotherapy, radiother	apy, surgery)	
0	l took today/ Y omitted today/N		
	d took today/ Y omitted today/N		
	or Blocker: Y and took today/ Y omitted today/N		
	took today/ Y omitted today/N ocker: Y and took today/ Y omitted today		
	today/Y omitted today/N		
Regular NSAIDs: Y/			
Regular PPI: Y/N			
Haemodynamics	a part C manthe at least 12h prior to the approxime re	am at rest.	
	e past 6 months, at least 12h prior to the operating ro c, Diastolic	uni, at rest.	
Heart r			
-	iately prior to induction of anaesthesia:		
Heart r	c, Diastolic rate		
		surgery) (we need to ask for units for each hospital)	
Creatinine			
Albumin			
Haemoglobin conce	entration		
SURGERY	tion / concer / cynlerater / fracture / bladin	a lothor	
	tion/cancer/exploratory/fracture/bleedin		
	ed in the eCRF from the sortsurgery.com	website):	
Details of type of sur	to favoured definitions, to slightly reduce	variability)	
Urgency	to lavoured demittions, to signify reduce	variability)	
• ·	V /N		
Cancer treatment NTRA-OPERATIVE	Y/N		
Start of anaesthesia:	hhmm DDMMYY		
Start of surgery:	hhmm DDMMYY		
End of surgery:	hhmm DDMMYY		
End of anaesthesia:	hhmm DDMMYY		
SURGICAL			
Estimated blood loss (EBI	L, ml): <250ml, 251-1000ml, 1001	-3000ml. >3000ml	
ANAESTHETIC	_,,	,	
Blood pressure			
•	ed blood pressure: Systolic/Diastolic (MAP	can be calculated)	
Anaesthesia: tick all appli			
	edation without securing airway/regional	/spinal/CSE/epidural	



SQUEEZE

Interventions:

Arterial line: Y/N Central venous line: Y/N

Intra-operative vasoactive drugs

	No	Y as bolus	Y as infusion
Angiotensin II			
Atropine			
Akrinor [®] (Cafedrin/Theodrenalin)			
Dobutamine			
Dopamine			
Ephedrine			
Epinephrine (Adrenaline)			
Etilefrine			
Glycopyrronnium			
Metaraminol			
Milrinone			
Nitrates			
Norepinephrine (Noradrenaline)			
Phenylephrine			
Vasopressin or terlipressin			
Other 1			

Was the patient receiving a vasopressor infusion prior to surgery starting: Y/N

Fluids and blood products received INTRA-operatively only, volume of

Crystalloid:

Colloid (starch, gelofusine, albumin): Packed red blood cells:

Fresh frozen plasma:

Platelets:

Whole blood or autotransfusion (in ml):

POST-OPERATIVE

EARLY EVENTS

- We are interested in which vasoactive drugs were given and how they were given.
- We have split all vasoactive drugs into those that are VASOPRESSORS (in green column) and those that are not (blue).
- We only want additional information (completion of CRF2) if it was POSTOPERATIVE, was a VASOPRESSOR and was INFUSED.

Vasoactive drugs					
Vasopressor	Not predominantly vasopressor				
Dopamine	Atropine				
Epinephrine (Adrenaline)	Dobutamine				
Metaraminol	Ephedrine				
Norepinephrine (Noradrenaline)	Etilefrine				
Phenylephrine	Glycopyrronnium				
Vasopressin or Terlipressin	Nitrates				
Akrinor ®	Milrinone				
Angiotensin II					
We appreciate that ma	any drugs have mixed actions				

Following the end of surgery, did the patient receive any

vasopressor boluses Y/N oral/enteral vasopressor (midodrine) Y/N

Y/N, if Y, then did this continue for more than 1 hour after the end of surgery: Y/N if yes then this fulfils our criteria for PVI, so please also complete CRF2.

Would you liked to have given some vasopressor infusion but lacked appropriate resources to permit this to occur safely?

LATE COMPLICATIONS = WITHIN FIRST WEEK

Organ support

infusion

Pulmonary Ventilation: invasive mechanical ventilation / NIV / both / neither Cardiovascular New dysrhythmia: AF/other/none Acute Myocardial Infarction (type 1, using WHO 4th universal definition) Renal Highest creatinine (within the first week) postoperatively: Value/Not available [we calculate KDIGO] Received renal replacement therapy: Y/N (excluding chronic RRT users) Gastrointestinal Received parenteral nutrition: Y/N



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Infection

Treated with antibiotics for a newly diagnosed infection: Y/N

If Y: skin or soft tissue / respiratory / urinary / abdominal / lines / other

Surgical

Accordion Severity Classification of Postoperative Complications (Annals 2009): 0 (none) to 4 (death)

END OF EPISODE (intra-hospital follow up to 30 days)

Did the patient receive PVI that started more than 24h following surgery?: Y/N During this admission, did the patient die: Y/N Date of discharge, death or end of observational period: DDMMYY

CRF2: Additional information for those who received postoperative vasopressor infusion (PVI)

PLEASE DO NOT complete if:

- receiving inotropes without vasopressors
- received vasopressor only intra-operatively or for less than one hour postoperatively
- received vasopressors starting more than 24 hours postoperatively

At one hour after the completion of surgery, is the patient:

Receiving continuous infusion of neuraxial anaesthesia/analgesia i.e. epidural infusion	Y/N
Still receiving a sedative infusion	Y/N
Still has an airway in place (endotracheal tube, tracheostomy or supraglottic airway)	Y/N

1. How was it initially assessed that this patient should receive a vasopressor infusion?

Options:

- 1. Already receiving a vasopressor infusion and attempts to lower the infusion rate produced unacceptable hypotension
- 2. It was decided that the patient would no longer benefit from further attempts to increase the cardiac output through administration of IV fluids and the blood pressure was unacceptably low. This was on the basis of:
 - A. clinical assessment alone (vital signs, examination, lab results)
 - B. clinical assessment AND a measurement of preload responsiveness using cardiac output monitoring (or some direct surrogate of)
 - C. clinical assessment AND a measurement of preload responsiveness using echocardiography
 - D. clinical assessment AND a previously established maximum for IV fluid administration has been met i.e. 2L or 20ml/kg etc...
 - E. other free text
 - F. unknown

Day 0 = the calendar day of the start of the operation

2. Organ failure scores

	Day 0	POD1	POD2	POD3	POD4	POD5	POD6
SOFA score							

3. Blood pressure target and levels

	Day 0	POD1	POD2	POD3	POD4	POD5	POD6
Target MAP (if known)							
Lowest recorded MAP							
Highest recorded MAP							

4. Vasoactive drug infusion details

	Day 0	POD1	POD2	POD3	POD4	POD5	POD6
Vasopressor infusion 1							
Vasopressor infusion 2							
Vasopressor infusion 3							



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Vasopressor infusion 4				
	 	 · · ·	 	

For each vasopressor drug, for each day, we want the highest infused rate – for example, noradrenaline 0.5 mcg/kg/min

	Day 0	POD1	POD2	POD3	POD4	POD5	POD6
Inotrope 1							
Inotrope 2							

For each inotropic drug, for each day, we want the highest infused rate - for example, milrinone 0.3 mcg/kg/min

5. Fluids

	Day 0	POD1	POD2	POD3	POD4	POD5	POD6
Fluid balance							

A value between -5000 and +20000, in millilitres. If <-2000 or >+5000 then eCRF to ask data entry person to double check. Represents the volume of fluid in (including medications, liquid feeding etc), minus the volume of fluid out (includes urine output, drain output etc...). Option not to provide this information if not known.

6. Organ support in the first 28 days

Total number of days of receipt of ventilation (invasive or NIV):

Total number of days of receipt of vasopressor infusion:

Total number of days of receipt of parenteral nutrition:

Total number of days of receipt of renal replacement therapy: