| Patient ID: CRF 1 | _ ⁻ | . | . - | | | | S | Sex: Mal | e / Fem |
|---|-----------------------------|-----------|--|------------|-------------|-----------|--------------|----------|------------|
| 0. Informed cor | isent | | | | | | | | |
| Is consent applicab | | | o No | | | If ye | s date of | DD-N | IMM-YY |
| Mandatory unless the | | explicit | | | | cons | sent: | | |
| and written exemptic | on from IRB | | o Yes | | | | | | |
| 1. Patient Infor | mation | | | | | | | | |
| 1.1. Year of | | | 1.2. | | | 1.3. | Height | | |
| Birth * | | | Weight | | | * | | | |
| | | | * | | | | | | |
| 1.4. Clinical Frailty | Scale * | | 1 Very fit | | \bigcirc | 6 Mc | derately Fr | rail | \bigcirc |
| | | | 2 Well | | \bigcirc | | verely Frail | | \bigcirc |
| | | | 3 Manag | - | \bigcirc | | ry severely | Frail | \bigcirc |
| | | | 4 Vulnera | | \bigcirc | | minally | | \bigcirc |
| | | | 5 Mildly | Frail | \bigcirc | 10 D | on't know | | \bigcirc |
| Previous medical | history * | | | | | | | | |
| 1.5. Coronary Arte | - | | ○ No | | | ΟYe | es | | |
| , 1.6. Cerebrovascu | | | ○ No | | | O Ye | | | |
| 1.7. Peripheral vas | | e | ○ No | | | O Ye | es | | |
| 1.8. Atrial fibrillati | | | ○ No | | | O Ye | es | | |
| 1.9. Heart failure | | | ○ No | | | | o Yes | | |
| 1.10. Hypertensio | n | | o No | | | | Yes | | |
| 1.11. Diabetes | | | ○ No O Insulin dependent O Non-insulin dep | | | lin depe | ndent | | |
| 1.12. Chronic liver | disease | | o No | | | 0 Ye | es | | |
| 1.13. Chronic resp | iratory disea | se | o No | | 0 C O | PD | 0 | Other | |
| 1.14. Long-term st | eroid use | | o No | | · | o Ye | es | | |
| 1.15. Regular med | lications (tic | k all tha | tannly lea | ve blank | if not a re | ogular m | edication) | | |
| ACE inhibitor | If \Box yes | | ok day of su | | o omitte | - | | O un | known |
| | \rightarrow | | | 0, | | | 0, | | |
| Alpha blocker | If \Box yes \rightarrow | o toc | ok day of su | irgery | o omitte | d day of | surgery | 0 un | known |
| Angiotensin | If □ yes | o too | ok day of su | irgery | o omitte | d day of | surgery | o un | known |
| receptor blocker | \rightarrow | | | | | | | | |
| Beta blocker | If □ yes | o too | ok day of su | irgery | o omitte | d day of | surgery | 0 un | known |
| | \rightarrow | | | | | | | | |
| Calcium channel | If 🗆 yes | o too | ok day of su | irgery | o omitte | d day of | surgery | O un | known |
| blocker | \rightarrow | | | | | | | | |
| Diuretic | If \Box yes \rightarrow | ○ toc | ok day of su | irgery | o omitte | d day of | surgery | 0 un | known |
| Regular NSAIDs | If \Box yes \rightarrow | o toc | ok day of su | irgery | o omitte | d day of | surgery | o un | known |
| Haemodynamics. | Leave blank | if not av | vailable | | | | | | |
| Measurements in | | | | 2h prior | to the ope | erating r | oom, at res | st. | |
| 1.16. Systolic: | - 1 | | 7 Diastolic | | | | Heart rate | | |
| Reading immediat | ely prior to i | | | | | | | | |
| 1.19. Systolic: | / | | 20 Diastolic | | | 1.21 | Heart rate | 2: | |
| Laboratory. Leave | blank if not | | | | nit | 1 | | | |
| 1.22. Creatinine: | | | , | | - | mg/c | ll or µmol/l | L | |
| 1.23. Albumin | | | | | | | g/L or µmo | | |
| | า | | | | | | g/L or mm | | |
| 1.24. Haemoglobii | | 1 | | | | | <u> </u> | | |
| 1.24. Haemoglobii | | | | | | | | | |
| Surgery Reason for sur | gery * | nfection | | \bigcirc | | | | | |





European Society of Anaesthesiology and Intensive Care

| Patient ID: - | - | | Sex: Male | / Female | |
|----------------------------------|------------------------------|------------|---------------------------|------------|--|
| Fra | acture 🔘 | | | | |
| Ble | eeding O | | | | |
| Ot | her 🔿 | | | | |
| 2.2 Surgical procedure | Breast | 0 | Orthopaedic | 0 | |
| * | Gynaecological | 0 | Plastics / Cutaneous | Ō | |
| (select single most appropriate) | Head and neck | Õ | Upper gastro-intestinal | Ō | |
| | Hepato-biliary | 0 | Neurological/spinal | Ō | |
| | Kidney / urological | Õ | Vascular | Ō | |
| | Lower gastro-intestinal | \bigcirc | Other | 0 | |
| 2.3 Severity | Minor | \bigcirc | | | |
| * | Intermediate | 0 | | | |
| | Major | Õ | | | |
| 2.4. ASA-PS: | ASA 1: Healthy person | | | \bigcirc | |
| * | ASA 2: Mild systemic disease | <u>.</u> | | 0 | |
| | ASA 3: Severe systemic disea | ise | | 0 | |
| | ASA 4 Severe systemic diseas | se that | t is a constant threat to | 0 | |
| | life. | | | | |
| | ASA 5 A moribund person wl | no is n | ot expected to survive | \bigcirc | |
| | without the operation. | | | | |
| 2.5. Urgency | Urgent | | | \bigcirc | |
| * | (includes emergency, expedi | ted, u | rgent and immediate) | | |
| | Not urgent | | | \bigcirc | |
| | (includes planned/elective) | | | | |

| 3. Operative | | | | | | | |
|--|----------------------------|-----------------|---------------|-------|--|--|--|
| 3.1. Date of anaesthesia induction * | DD-MMM-YY | | | | | | |
| 3.2. Time of anaesthesia induction * | HH:MM | | | | | | |
| 3.3. Date of end of surgery * | DD-MMM-YY | | | | | | |
| 3.4. Time of end of surgery * | HH:MM | | | | | | |
| 3.5. Estimated blood loss (mL) * | <250 | 251-1000 | 1001-3000 | >3000 | | | |
| | 0 | 0 | 0 | 0 | | | |
| 3.6 /3.7 Lowest intraoperative blood | Systolic: | | Diastolic: | | | | |
| pressure (paired) * | | | | | | | |
| 3.8. Anaesthesia: | | | Volatile | | | | |
| * | | | TIVA | | | | |
| (Tick all that apply) | Seda | | | | | | |
| | | Regional | | | | | |
| | | | | | | | |
| | | | Epidural | | | | |
| 3.9. Airway | | Endot | racheal tube | 0 | | | |
| * | | | lottic airway | 0 | | | |
| | O2 f | acemask or nasa | l cannula | 0 | | | |
| 3.10. Arterial line * | 0 No | | 0 Yes | | | | |
| 3.11. Central venous line * | 0 No | | 0 Yes | | | | |
| 3.12. Which Intra-operative vasoactive | Atropine | | | | | | |
| drugs | Akrinor [®] (Cafe | drin/Theodrenal | in) | | | | |
| | Dobutamine | | | | | | |
| [Tick all that apply] | Dopamine | | | | | | |
| | Ephedrine | | | | | | |
| | Epinephrine (A | drenaline) | | | | | |
| | Glycopyrronniu | ım | | | | | |
| | Metaraminol | | | | | | |





| Patient ID: - - - - | - | | | Sex: Male / Fe | emale |
|--|------------------|-------------------|-------|----------------------------|-------|
| | Milrinone | | | | |
| | Nitrates | | | | |
| | Norepinephrine | e (Noradrenaline) |) | | |
| | Phenylephrine | | | | |
| | Vasopressin or | Terlipressin | | | |
| 3.13. Was the patient receiving a vaso anaesthesia? * | pressor infusion | prior to | O NO | o Yes | |
| 3.14. Fluids and blood products received during surgery: | Crystalloid | | | | (mL) |
| | Colloid | | | | |
| | (starch-gelofus | sine-albumin) | | | (mL) |
| | Packed red blo | | | | · · · |
| | | | | | (mL) |
| | Fresh frozen p | lasma | | | |
| | | | | | (mL) |
| | Platelets | | | | |
| | | | | | (mL) |
| | Whole blood o | or | | | |
| | autotransfusio | 'n | | | (mL) |
| 4. Post-operative Following the end of surgery (within 2 | 4h): | | | | |
| 4.1. Did the patient receive enteral vas (i.e. MIDODRINE) * | sopressors? | ○ No | o Yes | | |
| 4.2. Did the patient receive boluses of * | vasopressors? | ○ No | o Yes | | |
| 4.3. Did the patient receive an infusion vasopressors? * | n of | ○ No (Stop her | - | (Continue wi and 4.3.2) | th |
| 4.3.1. If yes , did the infusion continut 1 hour from the end of surgery? | e or start after | 0 No | ○ Yes | | |
| 4.3.2. If Yes , Did this infusion start wit from the end of surgery? | hin 24 hours | ○ No | ○ Yes | | |
| | 4.3.1 and .4.3.2 | 2, complete CRF | 2 | | |

| 5. Outcomes | | | | | | | |
|-----------------------------|---------------------------------------|-------------------|-------------|---------------------|--|--|--|
| 5.1. Ventilation: | O No O Invasive mechanical | | | o Non Ir | Non Invasive Ventilation (NIV) | | |
| * | | ventilation (IMV) | | | | | |
| 5.2. Acute Myocardial Infa | arction * | o No | | 0 Yes | | | |
| 5.3. New onset atrial fibri | llation * | o No | | 0 Yes | | | |
| 5.4. New onset other dysr | hythmia * | o No | | 0 Yes | | | |
| 5.5. Renal: Highest creatir | nine | | | mg/dl o | r μmol/L | | |
| (within the first week) pos | | | indicate | indicate which unit | | | |
| 5.6. Renal replacement th | o No | | o Yes | o Yes | | | |
| 5.7. Parenteral nutrition * | : | 0 No | | o Yes | | | |
| 5.8. Antibiotics for a | o No | | o Yes (comp | olete 🖓) | o Unknown | | |
| newly diagnosed | 0 Skin or so | oft tissue o | | o Abdomir | Abdominal | | |
| infection * | Respirato | ry o | | o Lines | ວ Lines | | |
| | O Urinary | o Urinary | | | | | |
| 5.9. Accordion | ○ None | | | | | | |
| classification of surgical | Mild complication | | | | | | |
| complication | o Moderate | e complicatio | on | | | | |
| * | o Severe co | mplication | | | | | |





European Society of

Anaesthesiology and

Intensive Care

| Patient ID: | <u> _ - -</u> - | <u>· _ - - _</u> | Sex: Male / Female |
|-------------|------------------|--------------------|--------------------|
| | | 0 Dooth | |

| 0 De | O'Dealii | | | | | | |
|----------------------------------|----------|-----------|--|--|--|--|--|
| 5.10. Date of hospital discharge | e or | DD-MMM-YY | | | | | |
| date of intrahospital death: * | | | | | | | |
| 5.11. Stayed an inpatient for m | ore ONO | o Yes | | | | | |
| than 30 days? * | | | | | | | |

Clinical Frailty Scale*

I Very Fit – People who are robust, active, energetic and motivated. These people commonly exercise regularly. They are among the fittest for their age.

2 Well – People who have no active disease symptoms but are less fit than category I. Often, they exercise or are very active occasionally, e.g. seasonally.

3 Managing Well – People whose medical problems are well controlled, but are not regularly active beyond routine walking.

4 Vulnerable – While not dependent on others for daily help, often symptoms limit activities. A common complaint is being "slowed up", and/or being tired during the day.



5 Mildly Frail – These people often have more evident slowing, and need help in high order IADLs (finances, transportation, heavy housework, medications). Typically, mild frailty progressively impairs shopping and walking outside alone, meal preparation and housework.

6 Moderately Frail – People need help with all outside activities and with keeping house. Inside, they often have problems with stairs and need help with bathing and might need minimal assistance (cuing, standby) with dressing.



7 Severely Frail – Completely dependent for personal care, from whatever cause (physical or cognitive). Even so, they seem stable and not at high risk of dying (within ~ 6 months).

8 Very Severely Frail – Completely dependent, approaching the end of life. Typically, they could not recover even from a minor illness.



9. Terminally III - Approaching the end of life. This category applies to people with a **life expectancy** <6 months, who are not otherwise evidently frail.

Scoring frailty in people with dementia

The degree of frailty corresponds to the degree of dementia. Common **symptoms in mild dementia** include forgetting the details of a recent event, though still remembering the event itself, repeating the same question/story and social withdrawal.

In **moderate dementia**, recent memory is very impaired, even though they seemingly can remember their past life events well. They can do personal care with prompting.

In severe dementia, they cannot do personal care without help.

* I. Canadian Study on Health & Aging, Revised 2008.
2. K. Rockwood et al. A global clinical measure of fitness and frailty in elderly people. CMAJ 2005;173:489-495.

© 2007-2009. Version 1.2. All rights reserved. Geriatric Medicine Research, Dalhousie University, Halifax, Canada. Permission granted to copy for research and educational purposes only.



2.3. Severity of surgery:

<u>Minor</u>: Procedure < 30 minutes. Examples: arthroscopy without intervention, removal of cutaneous tumour, proctology procedures, biopsy or excision biopsy of small lesions, etc

<u>Intermediate</u>: Procedure performed in a dedicated operating room that may pose the risk of significant complications or tissue injury. Examples: laparoscopic cholecystectomy, arthroscopy with intervention, bilateral varicose vein removal, tonsillectomy, inguinal hernia repair, breast lump resection, haemorrhoidectomy, appendicectomy, partial thyroidectomy, cataract surgery, uvuloplasty, minimally invasive repair of vaginal prolapse, vaginal hysterectomy, fixation of mandibular fracture, etc

<u>Major</u>: Performed in a dedicated operating room and is expected to last more than 90 minutes. Examples: major gut resection, major joint replacement, mastectomy, extensive head and neck tumour resection, abdominal aortic aneurysm repair, major vascular bypass procedure, procedures involving free flap to repair tissue defect, amputation, total thyroidectomy, cystectomy, trans-urethral resection of prostate, resection of liver tumour, carotid endarterectomy, nephrectomy, total abdominal hysterectomy, spinal discectomy, etc





European Society of Anaesthesiology and Intensive Care

| CRF Z | | | | | | | |
|---|-----------------------------|-------|--|--|--|--|--|
| Postoperative vasopressor infusion | | | | | | | |
| 6.1. Did this patient have an infusion of vasopressors | 0 No | o Yes | | | | | |
| that was either started or continued at least 1 hour | (If 'No' then please do not | | | | | | |
| after surgery: * | complete any further) | | | | | | |

| At one hour after the completion of surgery, is the patient: | | | | | |
|--|------|-------|--|--|--|
| 6.2. Receiving continuous infusion of neuraxial O No O Yes | | | | | |
| anaesthesia/analgesia i.e. epidural infusion * | | | | | |
| 6.3. Still receiving a sedative infusion * | O NO | o Yes | | | |
| 6.4. Still has an airway in place (endotracheal tube, | O NO | o Yes | | | |
| tracheostomy or supraglottic airway) * | | | | | |

| Already receiving a vasopressor infusion and attempts to lower the infusion rate produced unacceptable hypotension, OR |
|---|
| • 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: |

| 7.1. SOFA score within 24 | (Use FAQ as required) |
|---------------------------|---|
| hours after surgery * | To calculate SOFA score: |
| [0-24] | https://clincalc.com/IcuMortality/SOFA.aspx |

| 7.2 - 7.8 MAP target (complete only if MAP is specified) | | | | | | | |
|---|--|--|--|--|--|--|--|
| Day 0 Day 1 Day 2 Day 3 Day 4 Day 5 Day 6 | | | | | | | |
| MAP | | | | | | | |

| 7.10 – 7.16 HIGHEST blood pressure for each day (paired) | | | | | | | | |
|---|---|--|--|--|--|--|--|--|
| On Postoperative unit/ICU only. Leave blank if not available. | | | | | | | | |
| | Day 0 Day 1 Day 2 Day 3 Day 4 Day 5 Day 6 | | | | | | | |
| Systolic | | | | | | | | |
| Diastolic Diastolic | | | | | | | | |





Patient ID: |_|_|-|_|-|_|_|-|__|

7.18 – 7.24 **LOWEST** blood pressure during the day (<u>paired</u>) On Postoperative unit/ICU only. Leave blank if not available.

| on rostoperative antifico only. Leave blank in not available. | | | | | | | |
|---|-------|-------|-------|-------|-------|-------|-------|
| | Day 0 | Day 1 | Day 2 | Day 3 | Day 4 | Day 5 | Day 6 |
| Systolic | | | | | | | |
| Diastolic | | | | | | | |

| 7.26 – 7.32 Vasoactive drug infusion, tick if applicable | | | | | | | |
|--|-------|-------|-------|-------|-------|-------|-------|
| | Day 0 | Day 1 | Day 2 | Day 3 | Day 4 | Day 5 | Day 6 |
| Noradrenaline | | | | | | | |
| Angiotensin II | | | | | | | |
| Dobutamine | | | | | | | |
| Dopamine | | | | | | | |
| Epinephrine (Adrenaline) | | | | | | | |
| Metaraminol | | | | | | | |
| Milrinone | | | | | | | |
| Phenylephrine | | | | | | | |
| Terlipressin | | | | | | | |
| Vasopressin | | | | | | | |

| Total number of days: * | | | | | |
|--|--|--|--|--|--|
| 7.33. receipt of ventilation (invasive or NIV) | | | | | |
| 7.34. receipt of vasopressor infusion | | | | | |
| 7.35. receipt of parenteral nutrition | | | | | |
| 7.36. receipt of renal replacement therapy | | | | | |
| 7.37. duration of stay in ICU/postoperative unit | | | | | |

| COVID | | | | | | | |
|---------------------------------------|------|------|----------------------|-------|-----------|--|--|
| 7.38. Did the patient | O NO | | o Yes | | o Unknown | | |
| have any testing for | | | If yes, answer below | | | | |
| SARS-CoV2? * | | | | | | | |
| 7.38.1. If Yes - did the patient test | | o No | | o Yes | | | |
| positive in the perioperative | | | | | | | |
| period? | | | | | | | |



