

Revision AAA Repair

Patient Details

Patient Consent* No Yes Not Required *If patient not consented:*
 Date consent recorded ____ / ____ / ____ (DD/MM/YYYY) *Do not record NHS number, name(s) or postcode.*
 NHS number* _____
 Date of birth* ____ / ____ / ____ (DD/MM/YYYY) *If consent not required:*
 Sex* Male Female *Ignore consent date.*
 Last name _____
 First name _____
 Postcode* _____

Admission Details

Admission date* ____ / ____ / ____ (DD/MM/YYYY)
 Mode of admission* Elective Non-elective
 Hospital code* *Pre-populated drop down menu on NVR audit site*
 Local ID* _____
 Procedure type **Abdominal Aortic Aneurysm Repair (Revision)**

Pre-operative: Pathway and Indications

AAA/Aortic Diameter* _____ in mm
 Previous Aortic op* No Open repair
 Endovascular repair AAA repair – type unknown
 Both open and endovascular repair
 Indication for re-intervention* AAA sac expansion
 Graft migration
 Graft stenosis or occlusion
 Please select as many options as possible Graft infection
 New arterial disease (proximal or distal to original arterial graft)
 Pseudo-aneurysm
 Other

- Pre-operative endoleak* No endoleak
 Type 1
 Type 2
 Type 3
 Type 4
 Type 5

Assessments (for Elective Pathway only)

CT/MR angio assessment date ___/___/_____ (DD/MM/YYYY)
MDT discussion date ___/___/_____ (DD/MM/YYYY)
Date of Anaesthetic review ___/___/_____ (DD/MM/YYYY)

Consultant vasc anaesthetist review
 No Yes

Fitness measurement used*

None
Please select as CPET
many options Incremental shuttle walk test/6 minute walk test
as applicable Non-invasive cardiac stress test
 Transthoracic echocardiogram

Investigation after preop anaesthetic assessment*

No additional investigation /intervention
Please select as Referral to another specialty
many options Optimisation/change in drug therapy
as possible Coronary angiogram

For both Elective and Non-elective pathways

Patient weight* _____ in Kg

Patient height* _____ in cm

Risk Scoring

- Comorbidities* None Chronic heart failure
 Diabetes Chronic renal disease
Please select as Hypertension Stroke
many options Chronic lung disease Active/managed cancer
as applicable. Ischaemic heart disease Lower limb arterial disease
- Smoking status* Current or stopped within 2 months Ex-smoker Never smoked

- White cell count* _____ (x10⁹/l)
Haemoglobin* _____ (dg/l)
Sodium* _____ (mmol/l)
Potassium* _____ (mmol/l)
Creatinine* _____ (µmol/l)
Albumin _____ (g/l)

Non Elective Admission – Additional items

- Highest pulse pre-op _____
Lowest systolic BP pre-op _____

- Abnormal ECG* Normal Abnormal
- ASA Grade* 1 – Normal
 2 – Mild disease
 3 – Severe, not life-threatening
 4 – Severe, life-threatening
 5 – Moribund patient

- Pre-operative medication* None Beta blocker
 Single anti-platelet ACE inhibitor / ARB s
 Dual anti-platelet Oral anti-coagulant
 Statin

- Peri-operative medication* None Antibiotic prophylaxis DVT prophylaxis

Has the patient had COVID-19 within the last 2 months? No Yes

COVID-19 Vaccine? No Yes, 1 dose Yes, 2 doses Yes, 3+ doses

- Patient's frailty score Not frail (well or managing well, routinely walking)
 Mild frailty (evident slowing such as difficulty walking outside)
 Moderate frailty (need help with some personal care or keeping house)
 Severe frailty (completely dependent for personal care)

Procedure: AAA Repair

Date/Time start ____ / ____ / _____ (DD/MM/YYYY); ____ : ____ (HH:MM)

- Aortic status*
- 1 Asymptomatic
 - 2 Symptomatic unruptured
 - 3 Ruptured
 - 4 Aortic transection
 - 5 Acute dissection
 - 6 Chronic dissection

Type of repair 4 Revision Open 5 Revision EVAR 3 Complex EVAR

OPCS code of procedure 1* _____ *All options will be available on NVR*

OPCS code of procedure 2 _____ *audit site via drop down menus.*

OPCS code of procedure 3 _____

- Re-intervention 1 Explant (partial or total) 6 Ligation of aortic branches
- Procedure(s)* 2 Extra-anatomical bypass 7 Embolisation
- 3 Relining 8 Proximal extension cuff / Fenestrated cuff
- 4 Endoanchors / Bare metal stent / Banding
- 5 Distal extension 9 Other

- Neck angle 1 0 to 60 degrees 3 75 to 90 degrees
- 2 60 to 75 degrees 4 More than 90 degrees

Neck diameter _____ in mm

Neck length _____ in mm

Open

- AAA Clamp site 1 Infra-renal 5 AAA Graft 1 Tube
- 2 Supra-renal 5 Bifurcated
- 3 Supra-mesenteric 6 Any groin incision
- 4 Supra-coeliac
- 5 Thoracic aorta

EVAR

Complex EVAR – Additional items

- EVAR exclusion* 0 No
- 1 Yes primarily
- 2 Yes after adjunctive procedures

- Extended External 0 No
- 1 Right

- Iliac Artery (EIA) 2 Left
- 3 Bilateral

Common Iliac Artery (CIA) diameter (larger one) _____ in mm

- Type of complex EVAR 1 FEVAR
- 2 BEVAR
- 3 TEVAR
- 4 Iliac branched graft
- 5 Composite graft
- 6 Chimney/periscope/snorkel

Note: If complex EVAR is 3=TEVAR, then the following questions won't appear on the screen: iliac branch, neck angle, neck length, extended into external iliac artery, common iliac artery diameter

- Iliac branch 0 No 1 Right 2 Left 3 Bilateral

* Mandatory fields

Perioperative Endoleak* <input type="checkbox"/> No endoleak <input type="checkbox"/> Type 1 <input type="checkbox"/> Type 2 <input type="checkbox"/> Type 3 <input type="checkbox"/> Type 4 <input type="checkbox"/> Unclassified <i>If NO perioperative endoleak ignore the next 2 items.</i> Endoleak intervention <input type="checkbox"/> No <input type="checkbox"/> Yes Success <input type="checkbox"/> No <input type="checkbox"/> Yes	
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Devices	
<i>Device section (please add each device component). The device barcode can also be scanned directly into the NVR IT system.</i>	
Manufacturer _____	Batch/Lot Number _____
Product Number _____	Was this device used on IFU? <input type="checkbox"/> No <input type="checkbox"/> Yes
GTIN _____	
Manufacturer _____	Batch/Lot Number _____
Product Number _____	Was this device used on IFU? <input type="checkbox"/> No <input type="checkbox"/> Yes
GTIN _____	

Operator	
Vascular specialist 1* _____	Anaesthetist 1* _____
Vascular specialist 2 _____	Anaesthetist 2 _____
Vascular specialist 3 _____	
Vascular specialist 4 _____	

Peri-operative Anaesthetic Details

- Anaesthetic type* Local infiltration
- Please select as many options as applicable* Plexus/compartiment block
- Neuraxial block (spinal/epidural)
- General anaesthetic
- Direct arterial monitoring No Yes
- Intraoperative cardiac output monitoring No Yes
- Postoperative coagulopathy No Yes
- Core temperature ≥ 36 °C at end of procedure No Yes
- Patient reported severe pain within 1 hour of surgery No Yes

Post Operative

- Destination after surgery* Ward Level 2 (HDU/PACU) Level 3 (ICU) Died in theatre
- Note: If Died in theatre is selected, the remaining questions in the post-operative section will not show*
- Critical care stay* _____ (Number of days)
- Return to theatre within admission* No Yes
- Readmission to a higher level of care* No Yes
- Postoperative complications* None
- Cardiac (MI / NSTEMI / heart failure)
- Respiratory
- Cerebral (stroke)
- Renal failure
- Please select as many options as applicable.* Haemorrhage
- Limb ischaemia
- Paraplegia
- Ischaemic bowel
- Post-operative confusion
- Major GI complication
- Surgical site infection
- Other

Discharge

Discharge status – Alive on discharge* No Yes

Date discharged/died* ____/____/____ (DD/MM/YYYY)

Discharge destination*
 Usual place of residence
 Rehabilitation
 Other hospital
 Intermediate care (e.g. nursing or care home)

Please also complete the [COVID-19 dataset, which can be found on our website](#).

Follow Up

Readmission to hospital within 30 days* No Yes

Was the readmission for vascular reasons? * No Yes

Did the patient die within 30 days of the procedure? * No Yes

Reason for NO follow up
 Died prior to planned follow-up after discharge
 Moved out of area
 Did not attend
 Other

Date clinic appointment attended ____/____/____ (DD/MM/YYYY) *(Only if follow up occurred)*

If you have any queries please contact us on 020 7869 6621 and nvr@rcseng.ac.uk