**Version Control**

**NELA Patient Audit Dataset**

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| **Version** | **Date** | **Changes** |
| 2.0 | 24/11/2014 | Changes made to dataset for 2nd year. |
| 2.1.1 | 02/04/2015 | Still in hospital at 60 days answer option added to question 7.7 |
| 2.1.2 | 02/07/2015 | Wording edited for question 2.9 |
| 3.1 | 01/12/2015 | Changes made to dataset for 3rdyear. |
| 3.1.1 | 21/03/2016 | Q1.9 wording edited |
| 4.1 | 01/12/2016 | Changes made to dataset for 4th year. |
| 4.1.1 | 21/12/2016 | Question 1.10b modified to includehospital transfers |
| 5.1 | 01/12/17 | Changes made to dataset for 5th year. |
| 6.1 | 01/12/18 | Changes made to dataset for 6th year. |
| 6.1.1 | 01/04/19 | Possum Calculation removed; Q3.2, 3.25, 6.2, 6.23,Q3.1, 6.1 Updated options |
| 7.1.1 | 01/12/19 | Changes made to dataset for Year 7. These should be used from December 1st 2019Removed: 1.10b, 1.11, 1.12, 2.1, 2.3, 2.4, 2.7a (combined with 2.7), 2.7b, 2.8b, 3.2, 3.5i, 5.3c,6.2, 7.4b, 7.9Wording changed: 3.1, 6.24, 7.3, New question 3.1a, 5.3, 6.17a |

This is the NELA proforma. All data entry will be carried out through an online data collection web tool. The web tool will be accessible via pc, tablets and mobiles

This audit is a continuous prospective audit with real time data collection. It is expected that clinical teams enter the data real time rather than retrospectively.

**On the NELA Webtool by default Quality Improvement (QI) questions are enabled. If you do not wish to collect data for one or more QI questions, the questions can be disabled. This is done on the NELA webtool.**

For queries, please contact info@nela.org.uk Web tool for data entry: <https://data.nela.org.uk/>

**This form is for information purposes only.**

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| **1.** | **Demographics and Admission** |  |
| **1.1** | NHS Number |  |
| **1.2** | Pseudo-anonymisation | Computer generated |
| **1.3** | Local patient id/hospital number |  |
| **1.4** | Date of birth |  |
|  | Age on arrival | *Age will automatically be calculated on web tool* |
| **1.5** | Sex | * Male / Female
 |
| **1.6** | Forename |  |
| **1.7** | Surname |  |
| **1.8** | Postcode |  |
| **1.9** | Date and time the patient first arrived at the hospital/Emergency department |  |
| **1.10** | What was the nature of this admission? | * Elective / Non-elective
 |
| **1.10b** |  No Longer Required |  |
| **1.11** |  No Longer Required |  |
| **1.12** |  No Longer Required |  |
| **1.13a** | Is this patient known to have a Learning Disability? | * Yes No  Unknown
 |
| **1.13b** | Is this patient known to have an Autistic SpectrumDisorder? | * Yes No  Unknown
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| **2** | **Pre-op****If the patient is returning to theatre as an emergency following previous elective****surgery, all answers should relate to the emergency laparotomy, not the previous elective surgery.** |
| **2.1** |  No Longer Required |  |
| **2.2** | Date and time that the decision was made to operate | Date (DD/MM/YYYY)* Date not known

Time (HH:MM)* Time not known
 |
|  | *If this is unavailable please enter date and time that this* |
|  | *patient was first booked for theatre for emergency* |
|  | *laparotomy* |
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| **2.3** | No Longer Required |  |
| **2.4** |  No Longer Required |  |
| **2.5** | No Longer Required |  |
| **2.6** | No Longer Required |  |
| **2.7** | Was an abdominal CT scan performed in the pre- operative period as part of the diagnostic work-up? If performed, how was this CT reported pre- operatively?*(If CT is reported by a registrar and validated by a consultant* ***before*** *surgery, select “in-house consultant”. If* ***not validated*** *by consultant before surgery, select**“registrar”)* | * Yes – reported by in-house consultant
* Yes – reported by in-house registrar
* Yes – reported by outsourced service
* Yes but NOT reported
* No CT performed
* Unknown
 |
| **2.7a** | No Longer Required |  |
| **2.7b** | No Longer Required |  |
| **2.7c** | Was there a discrepancy between the CT report and surgical findings that altered or delayed either thediagnosis or surgical management? | * Yes
* No
* Unknown
 |
| **2.8a** | No Longer Required |  |
| **2.8b** | No Longer Required |  |
| **2.9** | No Longer required |  |
| **2.10** | What was the date and time of the first dose of antibiotics following presentation to hospital? *(only relevant for non-elective admissions)* | * In theatre, or

Date (DD/MM/YYYY)* Date not known

Time (HH:MM)* Time not known
* Not Administered
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| **2.11a** | Was sepsis, with a NEWS2 >=5 or >=3 in any one variable or another diagnosis requiring urgent antibiotics e.g. peritonitis / perforation, suspected on admission? | * Yes
* No
* Unknown
 |
| **2.11b** | Was sepsis, with a NEWS2 >=5 or >=3 in any one variable and/or another diagnosis requiring urgent antibioticse.g. peritonitis / perforation, suspected at the time thedecision for surgery was made? | * Yes
* No
* Unknown
 |
| **2.12** | Using the Clinical Frailty Score (see help box), what was the patients pre-admission frailty status assessed as being? | * (1-3) - not frail
* 4 - vulnerable
* 5 - mildly frail
* 6 - moderately frail
* 7 - severely frail - completely dependent for personal care
* 8 - very severely frail
* 9 - Terminally ill
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| **3** | **Pre-op Risk stratification** |  |
| **3.1** | Prior to surgery, what was the risk of death for the patient that was entered into medical record?*For info, wording of relevant standard “An assessment of mortality risk should be made explicit to the patient and recorded clearly on the consent form and in the**medical record.”* | * Lower (<5%)
* High (>=5%)
* Not documented
 |
| **3.1a** | If documented, how was risk assessed?  | * Objective clinical score
* Clinical judgement
 |
| **3.1b** | If patient assessed to be high risk, which **consultants** were involved immediately preoperatively in the assessment, decision making process and care of this patient? This may be either direct or indirect care. *Please mark all that apply.* | * Consultant Surgeon
* Consultant Anaesthetist
* Consultant Intensivist
* None
 |
| **3.2** | No Longer Required |  |
| **3.3** | What was the **ASA** score? | * 1: No systemic disease
* 2: Mild systemic disease
* 3: Severe systemic disease, not life- threatening
* 4: Severe, life-threatening
* 5: Moribund patient
 |
| **3.4** | What was the most recent pre-operative value forserum Creatinine (micromol/l) |  |
|  | * Not performed
 |
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| **3.5** | What was the most recent pre-operative value for blood lactate – may be arterial or venous (mmol/l) |

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|  | * Not performed
 |

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| **3.5i** |  No Longer Required |  |
| **3.5ii** | What was the most recent pre-operative value for albumin (g/l)? |  | * Not performed
 |
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|  | **NELA Risk calculation** |  |
|  | **For questions, 3.6 to 3.22 please enter values closest to time of booking for theatre in order to calculate****NELA Risk score. Answers should reflect chronic *and* acute pathophysiology**. |

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| **3.6** | Serum Sodium concentration (mmol/l) |  |
| **3.7** | Serum Potassium concentration (mmol/l) |  |
| **3.8** | Serum Urea concentration (mmol/l) |  |
| **3.9** | Serum Haemoglobin concentration (g/dl) |  |
| **3.10** | Serum White cell count (x109 / l) |  |
| **3.11** | Pulse rate(bpm) |  |
| **3.12** | Systolic blood pressure (mmHg) |  |
| **3.13** | Glasgow coma scale |  |
| **3.14** | Select an option that best describes this patient’s **ECG** | * No abnormalities
* AF rate 60-90
* AF rate >90/ any other abnormal rhythm/paced rhythm/ >5VE/min/ Q, ST or T wave abnormalities
 |
| **3.15** | Select an option that best describes this patient’s**cardiac signs** and chest xray appearance | * No failure
* Diuretic, digoxin, antianginal or antihypertensive therapy
* Peripheral oedema, warfarin Therapy or CXR: borderline cardiomegaly
* Raised jugular venous pressure or

CXR: cardiomegaly |
| **3.16** | Select an option that best describes this patient’s**respiratory history** and chest xray appearance | * No dyspnoea
* Dyspnoea on exertion or CXR: mild COAD
* Dyspnoea limiting exertion to < 1 Flight or CXR: moderate COAD
* Dyspnoea at rest/rate > 30 at rest or CXR:

fibrosis or consolidation |
| 3.16a | No Longer Required |  |
|  | *Online web tool will automatically calculate Physiology severity score* |  |
| **3.17** | Select the **operative severity** of the intended surgical intervention (see help box for examples) | * Major
* Major+
 |
| **3.18** | Including this operation, how many operations has the patient had in the 30 day period prior to thisprocedure? | * 1
* 2
* >2
 |
| **3.19** | Based on your clinical experience of the intended surgery, please estimate the likely ***intra*operative blood loss** (ml) | * <100
* 101-500
* 501-999
* >=1000
 |
| **3.20** | Please select a value that best describes the likely degree of **peritoneal soiling** | * None
* Serous fluid
* Localised pus
* Free bowel content, pus or blood
 |
| **3.21** | What severity of malignancy is anticipated to be present? | * None
* Primary only
* Nodal metastases
* Distant metastases
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| **3.22** | Please select **urgency** of surgical intervention*(see help notes for additional information)* | * 3. Expedited (>18 hours)
* 2B. Urgent (6-18 hours)
* 2A. Urgent (2-6 hours)
* 1. Immediate (<2 hours)
 |
|  | *Online web tool will automatically calculate Operative severity score* |  |
| **3.23** | No Longer Required |  |
| **3.24** | No Longer Required |  |
| **3.25** | Not all investigations available for calculation of NELA Risk |  |
| **3.26** | Estimated mortality using NELA risk adjustment model*(Figure only provided if all data available)* | Calculated |  |  |
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| **4** | **Intra-op** |  |
| **4.1** | Date and time of entry in to operating theatre/anaesthetic room (not theatre suite) | Date (DD/MM/YYYY) Time (HH:MM)* Time not known
 |
| **4.2** | Senior surgeon grade*(this can include surgeon supervising in theatre but not necessarily scrubbed)* | * Consultant
* Post-CCT fellow
* SAS grade
* Research Fellow / Clinical Fellow
* Specialty trainee
* Other
 |
| **4.2a** | Consultant present/supervising: Name/GMC/specialty of operating or supervising consultant*(If consultant not present, enter name of supervising**consultant)* | (Please select consultant - Online) |
| **4.3** | Senior anaesthetist present in theatre | * Consultant
* Post-CCT fellow
* SAS grade
* Research Fellow / Clinical Fellow
* Specialty trainee
* Other
 |
| **4.3a** | Consultant present (or supervising) : Name/GMC of anaesthetist*(If consultant not present, enter name of supervising**consultant)* | (Please select consultant - Online) |
| **4.4** | How did you provide goal directed fluid therapy? | * Patient recruited to FLO-ELA trial \*
* Not provided
* Dynamic index e.g. Stroke volume, PPV, SVV
* Static index e.g. CVP
* Other, eg bioimpedence
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| **5** | **Procedure** |  |
| **5.1** | Is this the first surgical procedure of this admission? | * Yes- First surgical procedure after admission
* No - Surgery for complication of

previous elective general surgical procedure within the same admission* No – Previous 'non-abdominal/non-general surgical' procedure within same admission (eg previous hip replacement)
* Unknown
 |
| **5.2** | What is the indication for surgery?*(Please select all that apply)* | * Peritonitis
* Perforation
* Abdominal abscess
* Anastomotic leak
* Intestinal fistula
* Phlegmon
* Pneumoperitoneum
* Necrosis
* Sepsis
* Small bowel obstruction
* Large bowel obstruction
* Volvulus
* Internal hernia
* Pseudo-obstruction
* Intussusception
* Incarcerated hernia
* Obstructing incisional hernia
* Haemorrhage
* Hiatus Hernia/para-oesophageal hernia
* Ischaemia
* Colitis
* Abdominal wound dehiscence
* Abdominal compartment syndrome
* Acidosis
* Iatrogenic injury
* Foreign body
* Planned relook
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| **5.3.a** | Main procedure |  Abdominal wall closure following dehiscience Abdominal wall reconstruction Adhesiolysis Colectomy: left (including sigmoid colectomy and anterior resection) Colectomy: right (including ileocaecal resection) Colectomy: subtotal or panproctocolectomy Colorectal resection - other Debridement Defunctioning stoma via midline laparotomy Drainage of abscess/collection Enterotomy Evacuation of haematoma Exploratory/relook laparotomy only Gastrectomy: partial or total Gastric surgery - other Haemostasis Hartmann’s procedure Intestinal bypass Laparostomy formation Large incisional hernia repair with bowel resection Large incisional hernia repair with division of adhesions Peptic ulcer – oversew of bleed Peptic ulcer – suture or repair of perforation Reduction of volvulus Removal of foreign body Removal of gastric band Repair of intestinal fistula Repair of intestinal perforation Repair of para-oesophageal hernia Repair or revision of anastomosis Resection of Meckel’s diverticulum Resection of other intra-abdominal tumour(s) Revision of stoma via midline laparotomy Small bowel resection Splenectomy Stricturoplasty Washout only Not amenable to surgery |
| **5.3.b** | Second procedure (at same laparotomy) |

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| **5.3e** | Was a stoma formed (by any means)? | * Yes
* No
 |
| **5.4** | Procedure approach | * Open
* Laparoscopic
* Laparoscopic assisted
* Laparoscopic converted to open
 |
| **5.5** | Operative findings:*(Please select all that apply)**If unsure whether this patient is eligible for NELA please refer to help box* | * Abscess
* Anastomotic leak
* Perforation – peptic ulcer
* Perforation – small bowel/colonic
* Diverticulitis
* Intestinal fistula
* Adhesions
* Incarcerated hernia
* Volvulus
* Internal hernia
* Intussusception
* Stricture
* Pseudo-obstruction
* Gallstone ileus
* Meckel’s diverticulum
* Malignancy – localised
* Malignancy – disseminated
* Colorectal cancer
* Gastric cancer
* Haemorrhage – peptic ulcer
* Haemorrhage – intestinal
* Haemorrhage – postoperative
* Ulcerative colitis
* Other colitis
* Crohn's disease
* Abdominal compartment syndrome
* Intestinal ischaemia
* Necrotising fasciitis
* Foreign body
* Stoma complications
* Abdominal wound dehiscence
* Normal intra-abdominal findings
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| **5.6** | Please describe the peritoneal contamination present*(select all that apply)* | * None or reactive serous fluid only
* Free gas from perforation +/- minimal contamination
* Pus
* Bile
* Gastro-duodenal contents
* Small bowel contents
* Faeculent fluid
* Faeces
* Blood/haematoma
 |
| **5.7** | Please indicate if the contamination was; | * Localised to a single quadrant of the abdomen
* More extensive / generalised
 |

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| **6** | **Post-op Risk stratification** |  |
| **6.1** | At the end of surgery, what was the risk of death for the patient that was entered into medical record? | * Lower (<5%)
* High (>=5%)
* Not documented
 |
| **6.1a** | If documented, how was risk assessed?  | * Objective clinical score
* Clinical judgement
 |
| **6.2** | No Longer Required |  |
| **6.3** | Blood lactate – may be arterial or venous (mmol/l) | #* Not performed
 |
|  | **Post-operative NELA Risk calculation**Q 6.4 – 6.14 No Longer Required |  |
|  | Physiology severity score: |  |
| **6.15** | What was the operative severity? (see help box for examples) | * Major
* Major+
 |
| **6.16** | Including this operation, how many operations has the patient had in the 30 day period prior to this procedure? | * 1
* 2
* >2
 |
| **6.17** | Please select this patient’s measured/estimated intraoperative blood loss (ml) | * <100
* 101-500
* 501-1000
* >1000
 |
| **6.17a** | If the patient’s blood loss was greater than 500mls, was Tranexamic Acid given? | * Yes
* No
 |
| **6.18** | Please select the option that best describes this patient’s degree of peritoneal soiling | * None
* Serous fluid
* Local pus
* Free bowel content, pus or blood
 |
| **6.19** | What was the level of malignancy based on surgical findings | * None
* Primary only
* Nodal metastases
* Distant metastases
 |
| **6.20** | What was the NCEPOD urgency?*(see help notes for additional information)* | * 3. Expedited (>18 hours)
* 2B. Urgent (6-18 hours)
* 2A. Urgent (2-6 hours)
* 1. Immediate (<2 hours)
 |
|  | *Online web tool will automatically calculate Operative severity score* |  |

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| **6.21** | No Longer Required |  |
| **6.22** | No Longer Required |  |
| **6.23** | Not all investigations available for calculation of NELA Risk |  |
| **6.24** | Where did the patient go for continued post-operative care following surgery? | * Ward
* Critical Care *(includes Level 2 HDU or Level 3 ICU)*
* Extended recovery area within theatres (eg PACU or OIR)
* Enhanced care area on a normal ward
* Died prior to discharge from theatre complex
 |
| **6.24a** | At the end of surgery, was the decision made to placethe patient on an end of life pathway? | * Yes
* No
 |
| **6.25** | No Longer Required |  |
| **6.26** | Estimated mortality using NELA risk adjustment model*(Figure only provided if all data available)* | Calculated |

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| **7** | **Post-op – Some fields will need to be completed****on discharge or death** |  |
| **7.1** | Total length of post-operative critical care stay (rounded up to whole days).*Includes both ICU and HDU stay -see help box for additional information. Do not include LOS in PACU/other enhanced recovery area* | Number required |
| **7.2** | No Longer Required |  |
| **7.3** | For patients aged 65 or older, was the patient assessed by a consultant geriatrician during any part of the perioperative period? | * Yes
* No
* Unknown
 |
| **7.4** | Within this admission, did the patient have an **unplanned or planned** return to theatre in the post- operative period following their initial emergency laparotomy? | * Yes; unplanned return
* Yes; planned return
* Yes; unplanned AND planned return
* No
* Unknown
 |
| **7.4a** | What was the main indication for the **unplanned** return to theatre?*(Select most significant reason)* | * Anastomotic leak
* Abscess
* Bleeding or Haematoma
* Decompression of abdominal compartment syndrome
* Bowel obstruction
* Abdominal wall dehiscence
* Accidental damage to bowel or other organ
* Stoma viability or retraction
* Ischaemia/non-viable bowel
* Sepsis/inadequate source control
* Deteriorating patient
* Missed pathology at first laparotomy
* Other
* Unknown
 |

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| **7.4b** |  No Longer Required |  |
| **7.5** | Did the patient have an unplanned move **from the ward** to a higher level of care within 7 days of surgery? (do not include moves from HDU to ITU, or escalationfrom other enhanced area/PACU) | * Yes
* No
* Unknown
 |
| **7.6** | No Longer Required |  |
| **7.7** | Status at discharge | * Dead  Alive
* Still in hospital at 60 days
 |
| **7.8** | Date discharged from hospital | (DD/MM/YYYY)Date required |
| **7.9** | No Longer Required |  |
|  | **COVID-19 Questions** |  |
| **7.10** | Please indicate the patient's SARS-CoV-2/COVID-19 infection status |  Infected at time of surgery based on a recentpositive RT-PCR antigen (swab) testConsidered as infected at time of surgery onclinical grounds despite negative (ie false negative) or indeterminate antigen testPositive antigen test or clinical diagnosis ofCOVID-19 during admission but unable to determine whether pre/post-op from the medical recordNot infected at time of surgery based on clinical presentation AND negative swab but had a new positive antigen test or clinical diagnosis of COVID-19 post-operativelyConsidered to be not infected throughoutinpatient stayAntigen test not doneUnable to answer |
| **7.11** | Regardless of actual COVID status, was the patient managed as infected with COVID whilst in the theatre suite for their initial emergency laparotomy (this does not mean, was enhanced PPE used only for the AGPs) | YesNoUnable to answer |
| **7.12** | Please indicate the patient's SARS-CoV-2 antibody status | PositiveNegativeNot testedUnable to answer |