

Supplementary material



INSTITUTION

Important: This form must only include information based on routine practice at this **INSTITUTION**.

1. Type: (Please select one)

- Academic hospital
 Group practice
 Hospital
 Private
 Other (Please state)

2. Morning session time:

HH:MM To HH:MM

3. Afternoon session time:

HH:MM To HH:MM

4. Are patients informed about their procedure beforehand? (Please select one)

- Yes (Please select all that apply) No

- Patient information leaflet mailed before pre-procedure visit
 Patient information leaflet at pre-procedure visit
 Verbal information/Telephone call
 Other (Please state)

5. Are patients required to give their informed consent? (Please select one)

- Yes (signed)
 Yes (verbal)
 No

6. Are patients' blood pressures routinely monitored during and after procedure? (Please select one)

- Yes
 No
 Sometimes (Please state circumstances)

Supplementary material



7. Are patients' heart rates and oxygen saturation routinely measured during and after procedure? *(Please select one)*

- Yes
 No
 Sometimes *(Please state circumstances)*

8. Does a protocol exist for post-procedure care? *(Please select one)*

- Yes No

9. Who authorises patient discharge post-procedure? *(Please select one)*

- Endoscopist
 Qualified nurse
 Second doctor
 Other *(Please state)*

10. Is caecal intubation rate routinely recorded? *(Please select one)*

- Yes No

11. Is adenoma detection rate routinely recorded? *(Please select one)*

- Yes No

12. Is polyp detection rate routinely recorded? *(Please select one)*

- Yes No

13. Are CO₂ and irrigation devices regularly used? *(Please select one)*

- Yes No

14. Are during-procedure complications routinely recorded? *(Please select one)*

- Yes No

15. Are post-procedure complications routinely recorded? *(Please select one)*

- Yes No

Supplementary material



16. Is patient satisfaction recorded? (Please select one)

Yes (Please select all that apply) No

- Follow-up call
- Follow-up consultation
- Patient questionnaire
- Post-procedure evaluation
- Other (Please state)

17. Is scale-based bowel cleansing quality routinely recorded? (Please select one)

Yes (Please select one) No

- Aronchick Bowel Preparation Scale
- Boston Bowel Preparation Scale
- Chicago Bowel Preparation Scale
- Harefield Cleansing Scale
- Institution
- Ottawa Bowel Preparation Scale
- Other (Please state)

18. Are quality guidelines routinely followed? (Please select one)

Yes (Please answer the following) No

Type of guidelines? (Please select all that apply)

- International
- Local
- National

Guidelines:

Supplementary material



PRACTITIONER

Important: This form must only include information based on **YOUR** routine practice.

1. **Role: (Please select one)**

- Gastroenterologist
 Internist
 Nurse
 Surgeon
 Trainee

2. **How many total colonoscopies have you performed in the last 12 months? (Please select one)**

- Fewer than 300 300 or more

3. **Scope manipulation: (Please select one)**

- Practitioner only Practitioner and assistant

4. **Do you routinely use a cleansing scale? (Please select one)**

- Yes (Please select one) No

- Aronchick Bowel Preparation Scale
 Boston Bowel Preparation Scale
 Chicago Bowel Preparation Scale
 Harefield Cleansing Scale
 Ottawa Bowel Preparation Scale
 Personal
 Other (Please state)

5. **Do you routinely use a polyp classification scale? (Please select one)**

- Yes (Please select one) No

- Paris Classification Scale
 Personal
 Other (Please state)

Supplementary material



6. Do you routinely record caecal intubation rate? *(Please select one)*

Yes No

7. Do you routinely record adenoma detection rate? *(Please select one)*

Yes No

8. Do you routinely record polyp detection rate? *(Please select one)*

Yes No

9. Do you routinely record polyp removal rate? *(Please select one)*

Yes No

10. Do you routinely record polyp retrieval rate? *(Please select one)*

Yes No

11. Do you routinely place tattoos following polyp removal based on guidelines? *(Please select one)*

Yes No

12. Do you routinely record retraction time? *(Please select one)*

Yes No

Supplementary material



PROCEDURE

Patient Information

1. Year of birth:

 YYYY

2. Gender: *(Please select one)*

 Male Female

3. Height:

 cm

4. Weight:

 kg

5. BMI:

 kg/m²

6. Type: *(Please select one)*

 In-patient Out-patient

Pre-Procedure

7. Patient referred by: *(Please select one)*

- GP
- Own speciality
- Screening programme
- Self-referred
- Other *(Please state)*

Supplementary material



8. Reason for procedure: *(Please select one)*

Clinical signs and symptoms *(Please select all that apply)*

Altered bowel

Pain

Rectal bleeding

Other *(Please state)*

Follow-up

Following positive screening test

Previous unsuccessful procedure

Screening due to familial risk

Screening without pre-screening test

Other *(Please state)*

9. Bowel preparation used? *(Please select one)*

Yes *(Please answer the following)* No

Type, including additional products: *(Please select all that apply)*

2 litre PEG and ascorbate

4 litre PEG

Bisacodyl

Enema

Sodium phosphate

Sodium picosulphate and magnesium citrate

Tri-sulphate

Other *(Please state)*

Dosing regimen: *(Please select one)*

Evening

Same day

Split

Other *(Please state)*

Did the patient follow the bowel preparation instructions? *(Please select one)*

Yes No

Supplementary material



How much bowel preparation was consumed? *(Please select one)*

- 0 - 25%
- 25 - 50%
- 50 - 75%
- 75 - 99%
- 100%

How much fluid was consumed in total, including additional products? *(Please select one)*

- 0 - 1 litre
- 1 - 3 litres
- 3 - 6 litres
- Over 6 litres

Time period between last intake of bowel preparation and procedure:

hours

10. Time of colonoscopy:

HH:MM

11. Patient had a total colonoscopy in the last 5 years? *(Please select one)*

- Yes *(Please answer the following)* No

Date of last total colonoscopy:

MM YYYY

Procedure

12. Sedation-related medications administered? *(Please select one)*

- Yes *(Please select all that apply)* No

- Entonox
- General Anaesthesia
- Midazolam
- Opiates
- Propofol
- Other *(Please state)*

Supplementary material



13. Procedure-related medications administered? (Please select one)

Yes (Please select all that apply) No

- Buscopan
 Glucagon
 Other (Please state)

14. Medication administered by: (Please select all that apply)

- Anaesthetist
 Endoscopist
 Qualified nurse
 Second doctor
 Other (Please state)

15. Chromoendoscopy used? (Please select one)

Yes (Please select all that apply) No

- Localised (digital)
 Localised (dye)
 Pan (digital)
 Pan (dye)

16. High definition equipment used? (Please select one)

Yes (Please select all that apply) No Don't know

- Scope
 Monitor/Screen

17. Assistive technology used? (Please select one)

Yes (Please select all that apply) No

- Cap-assisted
 Endocuff-assisted
 Scope guide
 X-ray
 Other (Please state)

Supplementary material



18. Intended endpoint: (Please select one)

- Anastomosis
 Caecum
 Terminal ileum/Neo terminal ileum

19. Endpoint photo documented? (Please select one)

- Yes No

20. Intended endpoint reached? (Please select one)

- Yes No (Please select one)

- Insufficient preparation
 Pain
 Pathology encountered
 Stricture
 Technically difficult
 Unsafe procedure
 Other (Please state)

21. Retraction time recorded? (Please select one)

- Yes (Please state time) No

 minutes

22. Abnormal endoscopic finding? (Please select one)

- Yes (Please select all that apply) No

- Cancer
 Diverticulae
 Inflammation
 Polyps
 Other (Please state)

Supplementary material



Right Colon Segment Classification

23. Cleansing quality: *(Please select one)* *

- Excellent
 Good
 Poor
 Inadequate

24. Polyps detected? *(Please select one)*

- Yes *(Please answer the following)* No

Number of polyps:

Type of polyps: *(Please select all that apply)*

- Protruded lesion: Pedunculated polyp (Ip)
 Protruded lesion: Subpedunculated polyp (Isp)
 Protruded lesion: Sessile polyp (Is)
 Flat elevated lesion: Flat elevation of mucosa (o-IIa)
 Flat elevated lesion: Flat elevation with central depression (o-IIa/c)
 Flat lesion: Flat mucosal change (o-IIb)
 Flat lesion: Mucosal depression (o-IIc)
 Flat lesion: Mucosal depression with raised edge (o-IIc/IIa)

Size of largest polyp:

 mm

Were you able to characterise the endoscopic appearance of highest grade polyp? *(Please select one)*

- Yes *(Please select one)* No
- Adenoma
 Hyperplastic
 Malignant
 Sessile serrated

Supplementary material



Transverse Colon Segment Classification

25. Cleansing quality: *(Please select one)* *

- Excellent
 Good
 Poor
 Inadequate

26. Polyps detected? *(Please select one)*

- Yes *(Please answer the following)* No

Number of polyps:

Type of polyps: *(Please select all that apply)*

- Protruded lesion: Pedunculated polyp (Ip)
 Protruded lesion: Subpedunculated polyp (Isp)
 Protruded lesion: Sessile polyp (Is)
 Flat elevated lesion: Flat elevation of mucosa (o-IIa)
 Flat elevated lesion: Flat elevation with central depression (o-IIa/c)
 Flat lesion: Flat mucosal change (o-IIb)
 Flat lesion: Mucosal depression (o-IIc)
 Flat lesion: Mucosal depression with raised edge (o-IIc/IIa)

Size of largest polyp:

 mm

Were you able to characterise the endoscopic appearance of highest grade polyp? *(Please select one)*

- Yes *(Please select one)* No
- Adenoma
 Hyperplastic
 Malignant
 Sessile serrated

Supplementary material



Left Colon Segment Classification

27. Cleansing quality: *(Please select one)* *

- Excellent
 Good
 Poor
 Inadequate

28. Polyps detected? *(Please select one)*

- Yes *(Please answer the following)* No

Number of polyps:

Type of polyps: *(Please select all that apply)*

- Protruded lesion: Pedunculated polyp (Ip)
 Protruded lesion: Subpedunculated polyp (Isp)
 Protruded lesion: Sessile polyp (Is)
 Flat elevated lesion: Flat elevation of mucosa (o-IIa)
 Flat elevated lesion: Flat elevation with central depression (o-IIa/c)
 Flat lesion: Flat mucosal change (o-IIb)
 Flat lesion: Mucosal depression (o-IIc)
 Flat lesion: Mucosal depression with raised edge (o-IIc/IIa)

Size of largest polyp:

 mm

Were you able to characterise the endoscopic appearance of highest grade polyp? *(Please select one)*

- Yes *(Please select one)* No
- Adenoma
 Hyperplastic
 Malignant
 Sessile serrated

Supplementary material



*Boston Bowel Preparation Scale (BBPS)

Questions 23, 25, and 27 relate to the BBPS.

Excellent (3): Entire mucosa of colon segment seen well with no residual staining, small fragments of stool or opaque liquid.

Good (2): Minor amount of residual staining, small fragments of stool and/or opaque liquid, but mucosa of colon segment seen well.

Poor (1): Portion of mucosa of the colon segment seen, but other areas of the colon segment not well seen due to staining, residual stool and/or opaque liquid.

Inadequate (0): Unprepared colon segment with mucosa not seen due to solid stool that cannot be cleared.

The calculated total score and grade will be displayed following segment classification.

Procedure

29. Bowel cleansing considered acceptable for the purpose of the procedure? *(Please select one)*

Yes No

30. Endoscopic intervention? *(Please select one)*

Yes *(Please select all that apply)* No

- Dilation of stenosis
- Endoscopic mucosal resection
- Endoscopic submucosal dissection
- Haemostasis
- Perforation repair by endoscopist
- Polypectomy (complete)
- Polypectomy (incomplete)
- Stent insertion
- Tattooing
- Other *(Please state)*

Supplementary material



31. Immediate complications? *(Please select one)*

Yes *(Please select all that apply)* No

- Bleeding requires admission
 Perforation *(Please select one)*

- Repaired
 Not repaired

- Sedation-related
 Severe pain
 Other *(Please state)*

Post Procedure

32. Non-routine (immediate) repeat procedure required? *(Please select one)*

Yes *(Please select one)* No

- Further treatment or intervention required
 Insufficient bowel preparation
 Pain
 Pathology encountered
 Stricture
 Technically difficult
 Unsafe procedure
 Other *(Please state)*

33. Patient successfully discharged, if classified as out-patient? *(Please select one)*

Yes No

Evaluation

34. Time to complete this form:

minutes

Important: Adverse events should be reported using the AE reporting system in the relevant country.