

CLINICAL MONITORING AND SAFE DISCHARGE OF PATIENTS ATTENDING THE ENDOSCOPY UNIT GUIDELINE

This guidance does not override the individual responsibility of health professionals to make appropriate decision according to the circumstances of the individual patient in consultation with the patient and /or carer. Health care professionals must be prepared to justify any deviation from this guidance.

INTRODUCTION

Recovery documentation and discharge planning is integral to the discharge of patients recovering from endoscopic procedures. It is essential that those who have received intravenous conscious sedation return to normal conscious levels before discharge and that a written record of recovery observations and events are documented.

This guidance is provided to enable qualified staff to safely discharge patients attending the Endoscopy unit using documented criteria.

Lead Clinician(s)

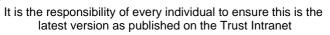
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Approved by SCSD Divisional Governance Meeting on: 5th April 2023

Review Date: 5th April 2026

This is the most current document and should be used until a revised version is in place

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline				
WAHT-MED-007 Page 1 of 11 Version 6				





Key amendments to this guideline

Date	Amendment	Ву:
12.07.06	Guideline approved on behalf of Clinical effectiveness	Dr Newrick
	Committee	Mrs Doherty
Nov 2008	Guideline approved on behalf of Clinical effectiveness	Mrs Doherty
	Committee	
02.08.10	Policy reviewed. Only change – updated references.	D Hathaway
11.05.12	Policy reviewed. Changes – Guideline 2.	D Hathaway
14.06.12	Policy reviewed. Changes to names of staff involved.	K Hinton / D Hathaway
12.07.12	Advised that document is not a policy, considered as	K. Hinton
	guidelines. Amended document accordingly.	
07.12.15	Guideline approved by Endoscopy directorate meeting	
25.6.2015	Addition of entonox and more specific guidance for	H Livett
	method of endoscopy	
November	Document extended whilst under review	TLG
2017		
December	Sentence added in at the request of the Coroner	
2017		
March	Document extended for 3 months as approved by TLG	TLG
2018		
June	Document extended for 3 months as approved by TLG	TLG
2018		
Jan 2019	Original author Karen Jeffries has since left organisation	L Mahachi
	and name removed from revised document.	
November	Guideline title amended to 'Clinical monitoring and safe	SCS Divisional
2019	discharge of patients attending the endoscopy unit guideline'	Governance Meeting
April 2023	Guideline reviewed and approved for 3 years	SCSD Divisional
		Governance Meeting

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit			
Guideline			
WAHT-MED-007 Page 2 of 11 Version 6			

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CLINICAL MONITORING AND SAFE DISCHARGE OF ALL PATIENTS ATTENDING ENDOSCOPY INCLUDING BOWEL CANCER SCREENING GUIDELINE

Introduction

Clinical Monitoring:

Endoscopic procedures can be undertaken with medication or without medication. Whether a patient had intravenous medication or not for the procedure, it is imperative that all patients' vital signs are monitored to enable a safe discharge. Academy of Medical Royal Colleges Safe Sedation Practice for Healthcare Procedures Standards and Guidance (2013) should be followed. (appendix 1)

Medication used during procedures or pre-existing comorbidities can contribute to patents deteriorating during endoscopy or during recovery. Benzodiazepines are common choice of sedative either alone or in combination with an opiate are usual in GI procedures. Such combination may increase the risk of oxygen desaturation and cardiorespiratory complications. There are also reported links between the use of local anaesthetic sprays and the development of pneumonia after gastroscopy. In addition, incidents have been reported locally where patient have suffered vasovagal during cystoscopy, hysteroscopy and sigmoidoscopy. It is therefore momentous that all patients are monitored as per table below.

Please note:

- 1) Safety and monitoring should be part of a quality assurance programme for endoscopy units.
- (2) Resuscitation equipment and drugs must be available in the endoscopy and recovery areas.
- (3) Staff of all grades and disciplines should be familiar with resuscitation methods and undergo periodic retraining.
- (4) Equipment and drugs necessary for the maintenance of airway, breathing, and circulation should be present in the endoscopy unit and recovery area (if outside the unit) and checked regularly.
- (5) A qualified nurse, trained in endoscopic techniques and adequately trained in resuscitation techniques, should monitor the patient's condition during procedures.
- (6) Before endoscopy, adverse risk factors should be identified. This may be aided by the use of a check list.
- (7) The dosage of all drugs should be kept to the minimum necessary. There is evidence that benzodiazepine/opioid mixtures are hazardous.
- (8) Specific antagonists for benzodiazepines and opioids exist and should be available in the event of emergency.
- (9) A cannula should be placed in a vein during endoscopy on 'at risk' patients.
- (10) Oxygen enriched air should be given to 'at risk' patients undergoing endoscopic procedures.
- (11) The endoscopist should ensure the well-being and clinical observation of the patient undergoing endoscopy in conjunction with another individual. This individual should be a qualified nurse trained in endoscopic techniques or another medically qualified practitioner.
- (12) Monitoring techniques such as pulse oximetry are recommended.
- (13) Clinical monitoring of the patient must be continued into the recovery area.
- (14) Records of management and outcome should be collected and will provide data for appropriate audit (Bell1991).

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit			
Guideline			
WAHT-MED-007 Page 3 of 11 Version 6			

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Requirements for clinical monitoring of endoscopy patients

(Deteriorating patients should be monitored using the NEWS chart and intervene as per accompanying action table)

Procedure All observations should now be documented on a NEWS 2 chart ready for escalation if needed. Full set of obs includes temp to enable accurate NEWS 2 score	On admissio n	Arrival in procedure room	After sedation	Midway eg sphincteromy /caecum NB: BP can be done if any concerns	End of procedure	Recovery full set of observations (obs) every 15 minutes total of 3-4 times depending upon individual patients following sedation For all other patients
Colonoscopy- No sedation or with Entonox only				Concerns		 ✓ - Once only unless therapeutic/EMR need to observe for signs of bleeding
Colonoscopy Sedation Gastroscopy- throat spray (inc GIB)						✓ One set post procedure only. If done in room no need for further set.
Gastroscopy – Sedation (inc GIB)						✓
EUS with Sedation						✓
Sigmoidoscopy- No sedation or Entonox						 ✓ - Once only unless therapeutic/EMR need to observe for signs of bleeding
Sigmoidoscopy- Sedation						✓
ERCP						✓ The recovery phase should be guided by the patients clinical condition and observations. But it is expected a minimum recovery time of 90 minutes to ensure the patient is adequately fit for discharge.
Bronchoscopy / EBUS with Sedation						✓
Cystoscopy	+ urinalysis					х

KEY

Full set Observations

Pulse & SPO2 only

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline			
WAHT-MED-007 Page 4 of 11 Version 6			

WAHT-MED-007

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NOTE: All patients should be assessed on an individual basis, monitoring of BP should be dictated by co-morbidities, all decisions should be clearly documented in the care document. Initial full set of obs should include temperature to accurately complete a NEWS score. Afterwards a temperature may not be required unless deterioration of the patient whereby NEWS needs to be calculated at every set of obs. ERCP should have temperature in recovery phase every time to calculate NEWS score and escalate where appropriate.

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline			
WAHT-MED-007 Page 5 of 11 Version 6			

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Discharge planning:

Recovery documentation and discharge planning is integral to the discharge of patients recovering from endoscopic procedures. It is essential that those who have any procedure with or without intravenous conscious sedation, opiates or Entonox return to normal preadmission consciousness and physical state before discharge. Documentation of recovery observations and events must be completed and signed prior to discharge.

Guidelines

A registered nurse trained within endoscopy will complete the discharge check list prior to the arrangement for patients going home or ward. All Bowel cancer screening patients will be seen by a Screening Nurse Practitioner prior to final discharge.

All adverse events should be discussed with the clinician in charge of that list and the nurse in charge of the unit and datixed as per trust policy and as guided by the list of endoscopy adverse events

Patient must have mental capacity or have an accompanying adult with them to receive specific discharge information and accompany them home.

If the patient does not wish to stay in endoscopy for the suggested length of recovery time, then the patient must sign a self-discharge form which is available from CLIP. (appendix 2)

If there is a suspicion that the patient is driving themselves home following sedation, then this should be discussed with the patient and advice give as per trust policy and a self-discharge form signed accordingly. In the case of a patient absconding before formal discharge then clear documentation in the nursing care pathway should be maintained.

In the case of any safeguarding concerns please refer to the intranet. Go to departments a-z and select safeguarding. Please follow the guidelines and links set out on that page. For reference, the contact details for the Integrated Safeguarding Team are:

Phone: 01905 733871 Ext: 33735

Email: wah-tr.SafeguardingWorcsacute@nhs.net

Office Hours: 09:00-16:30

In the case of emergency outside of office hours please follow the links on the intranet for emergency referrals for adults and children

Safeguarding children flowchart (appendix 3)

Safeguarding adult factsheet (appendix 4)

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline			
WAHT-MED-007	Page 6 of 11	Version 6	



Checks prior to discharge

Lignocaine used for upper gastrointestinal endoscopy

- Discharge from the room or recovery area as appropriate
- Stable vital signs
- Pain free or pain score as prior to admission
- Nausea free
- All appropriate information given to patient verbally and written; including nil by mouth for 30 minutes' post procedure

Follow up, from procedure is documented on the report, nursing documentation and patient information, unless consultant deems unsuitable.

Lignocaine and sedation utilised for upper GI procedures

- Patient is transferred to the recovery area complete with handover, monitoring and oxygen provision as per trust guidance
- Discharge as per Sedation guidance below
- Fluids and diet should not be offered for 30 minutes' post procedure.

Sedation and/or opiates only:

- Transferred to recovery area prior to discharge
- Monitoring post procedure (including BP and pulse oximetry)
- Stable vital signs
- Pain free or pain score as prior to admission
- Nausea free
- Alert and orientated as prior to admission
- Tolerating diet and fluid
- Cannula removed and documented post diet and fluid
- Suitable transport home.
- Responsible adult to stay with patient for 12 hours

Follow up from procedure is documented on the report, nursing documentation and patient information, unless consultant deems unsuitable.

Entonox then discharge should be to the recovery room

- Stable vital signs.
- Pain free or pain score as prior to admission
- Nausea free
- Alert and orientated as prior to admission

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline			
WAHT-MED-007 Page 7 of 11 Version 6			

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Follow up from procedure is documented on the report, nursing documentation and patient information, unless consultant deems unsuitable.

• The registered nurse will document, sign and date the discharge check list. Any deviation from the above must be discussed with the appropriate medical staff and fully documented.

Monitoring Tool

STANDARDS

- Accurate documentation of discharge will be audited twice yearly and discussed at unit meetings. Clinical Governance aware of results and action plan completed according to need
- 2. Datix and adverse events are monitored monthly at the Directorate Governance Meeting. Clinical Governance aware of results and action plan completed according to need.
- 3. Annual patient survey completed with a focus on patient aftercare. Discussed within each unit and at EUG, Clinical Governance and Patient Participation Forum (PPF)

References

- Academy of Medical Royal colleges (2013) Safe Sedation Practice for Healthcare Procedures Standards and Guidance. The Academy of Medical Royal Colleges, London.
- Baldwin.T (2010) "Policy of nurse led discharge of patients attending the Intervention Suite Kidderminster"
- Barson.A (2005) "Conscious sedation guidelines for medical and non-medical staff working within endoscopy units"
- British Society of gastroenterology (2003) "Guidelines on safety and Sedation during endoscopic procedures.
- Bell, G D (1991) Recommendations for standards of sedation and patient monitoring during gastrointestinal endoscopy. Gut, 32,823-827 British Society of Gastroenterology M A Quine, G D Bell, R F McCloy, J E Charlton, H B Devlin, A
- Hopkins (1995) Prospective audit of upper gastrointestinal endoscopy in two regions of England: safety, staffing, and sedation methods. Gut; 36: 462-467 British Society of Gastroenterology
- Joint Advisory Group on Gastrointestinal Endoscopy (JAG) accreditation standards for endoscopy services (2016)
- Somchai Amornyotin (2013) Sedation and monitoring for gastrointestinal endoscopy. World J Gastrointest Endosc. 2013 Feb 16; 5(2): 47–55. Published online 2013 Feb 16. doi: 10.4253/wjge.v5.i2.47
- Dunkley I et al (2018) UK consensus on non-medical staffing required to deliver safe, quality-assured care for adult patients undergoing gastrointestinal endoscopy. Frontline Gastroenterology 2018;0:1–11

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline			
WAHT-MED-007 Page 8 of 11 Version 6			



Contribution List

Key individuals involved in developing the document

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Karen Jefferies	JAG Lead/Clinical Governance
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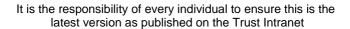
Circulated to the following individuals for comments

Name	Designation
Members of the Endoscopy Directorate	
Dr James Hutchinson	Consultant Anaesthetist

Appendix

Appendices:	
Appendix 1	Safe_Sedation_Practic e_1213.pdf
Appendix 2	Instructions to print self-discharge from C
Appendix 3	Referral to CSC Flow Diagram (1).docx
Appendix 4	Safeguarding Adult factsheet.pdf

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit		
Guideline		
WAHT-MED-007	Page 9 of 11	Version 6





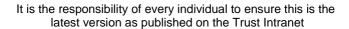
Supporting Document 1 - Equality Impact Assessment Tool

To be completed by the key document author and attached to key document when submitted to the appropriate committee for consideration and approval.

		Yes/No	Comments
1.	Does the policy/guidance affect one group less or more favourably than another on the basis of:		
	Race	No	
	Ethnic origins (including gypsies and travellers)	No	
	Nationality	No	
	Gender	No	
	Transgender	No	
	Religion or belief	No	
	Sexual orientation including lesbian, gay and bisexual people	No	
	Age	No	
	Disability - learning disabilities, physical disability, sensory impairment & mental health problems	No	
2.	Is there any evidence that some groups are affected differently?	No	
3.	If you have identified potential discrimination, are any exceptions valid, legal and/or justifiable?	Yes	
4.	Is the impact of the policy/guidance likely to be negative?	No	
5.	If so can the impact be avoided?		
6.	What alternatives are there to achieving the policy/guidance without the impact?		
7.	Can we reduce the impact by taking different action?		

If you have identified a potential discriminatory impact of this key document, please refer it to Human Resources, together with any suggestions as to the action required to avoid/reduce this impact. For advice in respect of answering the above questions, please contact Human Resources.

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline		
WAHT-MED-007 Page 10 of 11 Version 6		Version 6





Supporting Document 2 – Financial Impact Assessment

To be completed by the key document author and attached to key document when submitted to the appropriate committee for consideration and approval.

	Title of document:	Yes/No
1.	Does the implementation of this document require any additional Capital resources	No
2.	Does the implementation of this document require additional revenue	No
3.	Does the implementation of this document require additional manpower	No
4.	Does the implementation of this document release any manpower costs through a change in practice	No
5.	Are there additional staff training costs associated with implementing this document which cannot be delivered through current training programmes or allocated training times for staff	No
	Other comments:	

If the response to any of the above is yes, please complete a business case and which is signed by your Finance Manager and Directorate Manager for consideration by the Accountable Director before progressing to the relevant committee for approval

Clinical Monitoring and Safe Discharge of Patients Attending the Endoscopy Unit Guideline		
WAHT-MED-007	Page 11 of 11	Version 6