The Management of Constipation in Adult Patients Receiving Palliative Care

Clinical Audit Tool

2015



This clinical audit tool accompanies the clinical guideline: 'The Management of Constipation in Adult Patients Receiving Palliative Care' Issue date: 2015

This tool is a support tool for clinical audit based on the NCEC guideline. It is not NCEC guidance.

The audit could be carried out in any service where specialist or non-specialist healthcare professionals prescribe medications for the management of cancer pain. For example, GP practices, pharmacies and

The audit should involve clinical and non-clinical stakeholders, which may include medical staff of all grades, nurses, GPs, pharmacists, clinical audit staff and patients. Further information about patient and public involvement in clinical audit is available on the HSE website.

The audit standards are based on the National Clinical Effectiveness Committee guideline for the management of constipation in adult patients receiving palliative care. In developing this tool consideration has been given to the clinical issues covered by the guideline and the potential challenges of data collection. There may be other recommendations within the guideline suitable for the development of audit

To ask a question about this clinical audit tool, or to provide feedback to help inform the development of future tools, please email the National Clinical Programme for Palliative Care at

Recommendation	Guidance reference
ASSESSMENT	
	1.1
1. A thorough history and physical examination are recommended	
as essential components of the assessment process.	
See data collection form question a, b and c	
2. A digital rectal examination (DRE) should be considered to	1.3
exclude faecal impaction if it has been more than 3 days since the	
last bowel movement or if the patient complains of incomplete	
evacuation (following appropriate DRE training).	
See data collection form question d	
3. A plain film of the abdomen (PFA) is not recommended for routine	1.5
evaluation but may be useful in combination with history and	
examination in certain patients.	
See data collection form question e	
PREVENTION	
4. Education on the importance of pharmacological and non-drug	2.1
measures is essential to enable patients and caregivers to take an	
active role in constipation prevention.	
See data collection form question f	
NON PHARMACOLOGICAL MANAGEMENT	
5a. Attention should be paid to the provision of optimised toileting	3.1
while ensuring adequate privacy and dignity for all patients.	0.1
5b. Consideration should be given to lifestyle modification including	3.2
the adjustment of diet and activity levels within a patient's limitations.	0.2
See data collection form question g	
PHARMACOLOGICAL MANAGEMENT	
6. The combination of a softening and a stimulating laxative is often	4.3
required. Optimisation of a single laxative is recommended prior to	_
the addition of a second agent.	
See data collection form question h and i	
7. The laxative dose should be titrated daily or alternate days	4.4
according to response.	
See data collection form question j	
OPIOID INDUCED CONSTIPATION	
8. The development of opioid induced constipation should be	5.1
anticipated. A bowel regimen should be initiated at the	
commencement of opioid therapy.	
See data collection form question k	
9. In the management of opioid induced constipation, optimised	5.2
monotherapy with a stimulant laxative is essential followed by the	
addition of a softener if required.	
See data collection form question I	
10. The use of opioid receptor antagonists under specialist guidance	5.4
should be considered in patients whose treatment is resistant to	
conventional laxative therapy.	

See data collection form question m	
INTESTINAL OBSTRUCTION	
11. A stool softener should be considered in partial intestinal obstruction. Stimulant laxatives should be avoided.	6.1
See data collection form questions n and o	
12. In complete intestinal obstruction, the use of all laxatives should be avoided as even softening laxatives have some peristaltic action.	6.2
See data collection form question p	

Exceptions	Definitions
Patients who are actively dying	
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Patients who decline this procedure;	
patients with a stoma; patients with	
prostatic abscesses or prostatitis.	
Caution is advised when considering	
a DRE in immuno-compromised or	
thrombocytopaenic patients.	
Patients with reduced level of	None
consciousness; education should be	
tailored to the needs of individuals	
with cognitive impairment.	
None	None
None	
None	
Patients with stomas.	
I	

Audit Data for 'The Management of Constipation in Adult Patients F

			Question a	Question b
				Was a thorough physical
			Was an appropriate bowel history taken on initial	examination conducted?
			assessment?	examination conducted:
			assessment	
Audit ID	Age	Sex		
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Yes	0
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Total	0
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Demographics

Age range:	0 - 0
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Male	0
Female	0

Receiving Palliative Care' clinical audit

Question c	Question d	
If constipation was	Was a digital rectal examination (DRE	
identified, were the	appropriate DRE traing) to exclude faecal impaction in the following	
following components of a	groups of patients:	
comprehensive		
assessment completed:		
onset of symptoms,		
aggravating and alleviating		
factors, frequency and		
pattern of bowel motions,	Patients in whom it has been more than 3	If the patient complains of
stool volume and	days since the last bowel movement?	incomplete evacuation?
appearance, nausea,		
abdominal discomfort,		
bloating, flatus, tenesmus		
bloating, natus, terresinus		
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Ques	Question f	
Was a plain film of		Was education provided on
abdomen performed to	basis of specific	bowel management
assess for constipation?	consideration rather than	strategies to enable
	being done "routinely" e.g.	patients and caregivers to
	unreliable history,	take an active role in
	possibility of overflow	constipation prevention?
	diarrhoea?	
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Question g	Question h	Question i
Were non-pharmacological	In patients in whom more	Was optimisation of a
	than one laxative was used,	single laxative achieved
constipation management	was a combination of a	prior to the addition of a
plan (e.g. optimized	softening and a stimulating	second agent?
toileting, diet and lifestyle	laxative used?	
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laxative dose titrated dailyinitiated at thestimulant laxative achievedor alternate days accordingcommencement of opioidprior to the addition of a			
or alternate days according commencement of opioid prior to the addition of a	where required, was the		was optimisation of a
to response? therapy? softening laxative? softening laxative?			
	to response?	therapy?	softening laxative?
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Ques	Question n	
Did the patient demonstrate resistance to conventional	opioid receptor antagonist	In patients with partial
laxative therapy?	(e.g. methylnaltrexone, naloxegol, naloxone	
	containing preparation)	
	considered under specialist guidance?	
	v	Was the use of a stool
		softener considered?

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Question o	Question p
intestinal obstruction:	In patients with complete intestinal obstruction, was the use of all laxatives avoided?
	avoided?
Were stimulant laxatives avoided?	

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			Question a	Question b
			Was an appropriate bowel	Was a thorough physical
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			assessment?	examination conducted.
			assessment	
Audit ID	Age	Sex		
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Yes	0
Νο	0
Total	0
Percentage	#DIV/0!

Demographics

Age range:	0 - 0
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Male	0
Female	0

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stool volume and	days since the last bowel movement?	incomplete evacuation?
appearance, nausea,		
abdominal discomfort,		
bloating, flatus, tenesmus		
bloating, natus, terresinus		
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	on e	Question f
		Was education provided on
abdomen performed to	basis of specific	bowel management
assess for constipation?		strategies to enable
		patients and caregivers to
		take an active role in
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		constipation prevention?
	diarrhoea?	
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Question g	Question h	Question i
Were non-pharmacological	In patients in whom more	Was optimisation of a
	than one laxative was used,	single laxative achieved
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plan (e.g. optimized	softening and a stimulating	second agent?
toileting, diet and lifestyle	laxative used?	
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Question j	Question k	Question I
Where required, was the	Was a bowel regimen	Was optimisation of a
laxative dose titrated daily	initiated at the	stimulant laxative achieved
or alternate days according	commencement of opioid	prior to the addition of a
to response?	therapy?	softening laxative?
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Ques	tion m	Question n
Did the patient demonstrate		In patients with partial
resistance to conventional	opioid receptor antagonist	
laxative therapy?	(e.g. methylnaltrexone,	
	naloxegol, naloxone	
	containing preparation)	
	considered under specialist	
	guidance?	
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		Was the use of a stool
		softener considered?
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Question o	Question p
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