

## Guidance on the Management of Adult Patients with a High Output Stoma

This guidance covers the management of adult patients over 18 years old with a high output ileostomy, jejunostomy or colostomy.

High output stoma (HOS) management can be challenging and requires a multidisciplinary approach to aid management as each patient is different. Diagnosis, remaining bowel length/function will vary in each patient.

Stoma output volumes greater than 1500ml in 24 hours for more than 3 days should be treated as a high output.

- **AIM:** stoma output approximately 1000ml **and** able to maintain own hydration / fluid and electrolyte balance with PO intake.
- **Newly formed stoma** – seek advice from senior clinician / surgeon. Usually after surgery the stoma output volume begins to settle. However, this adaptation in bowel function occurs over days to months and is individual for every patient.

**A** INITIAL 48 hour management of HOS →

If high output persists 3 days after introduction of diet then follow pathway A below.

- **Bowel length** - Knowledge of remaining bowel length from duodeno-jejunal flexure (DJ flexure) and position of stoma is **essential in the management of HOS**.
- In patients **with short bowel syndrome** (bowel length less than 200cm from DJ flexure or continued high output despite conservative measures), continued dietetic input will be needed to meet enteral requirements and fibre is unlikely to be tolerated.

**A** INITIAL 48 hour management of HOS →

Start the management strategies in pathway A below.

Please note: these patients will need escalation to pathway B very quickly.

→ **B** CONTINUED management of HOS:

There may be a need for referral to a tertiary intestinal failure specialist centre for long-term management and long-term parenteral nutrition in short bowel patients. Please ensure a referral is made to the Nutrition Support Team to facilitate this.

**Nutrition Support e-REFER:** <https://intranet.gloshospitals.nhs.uk/departments/corporate-division/erefer/>

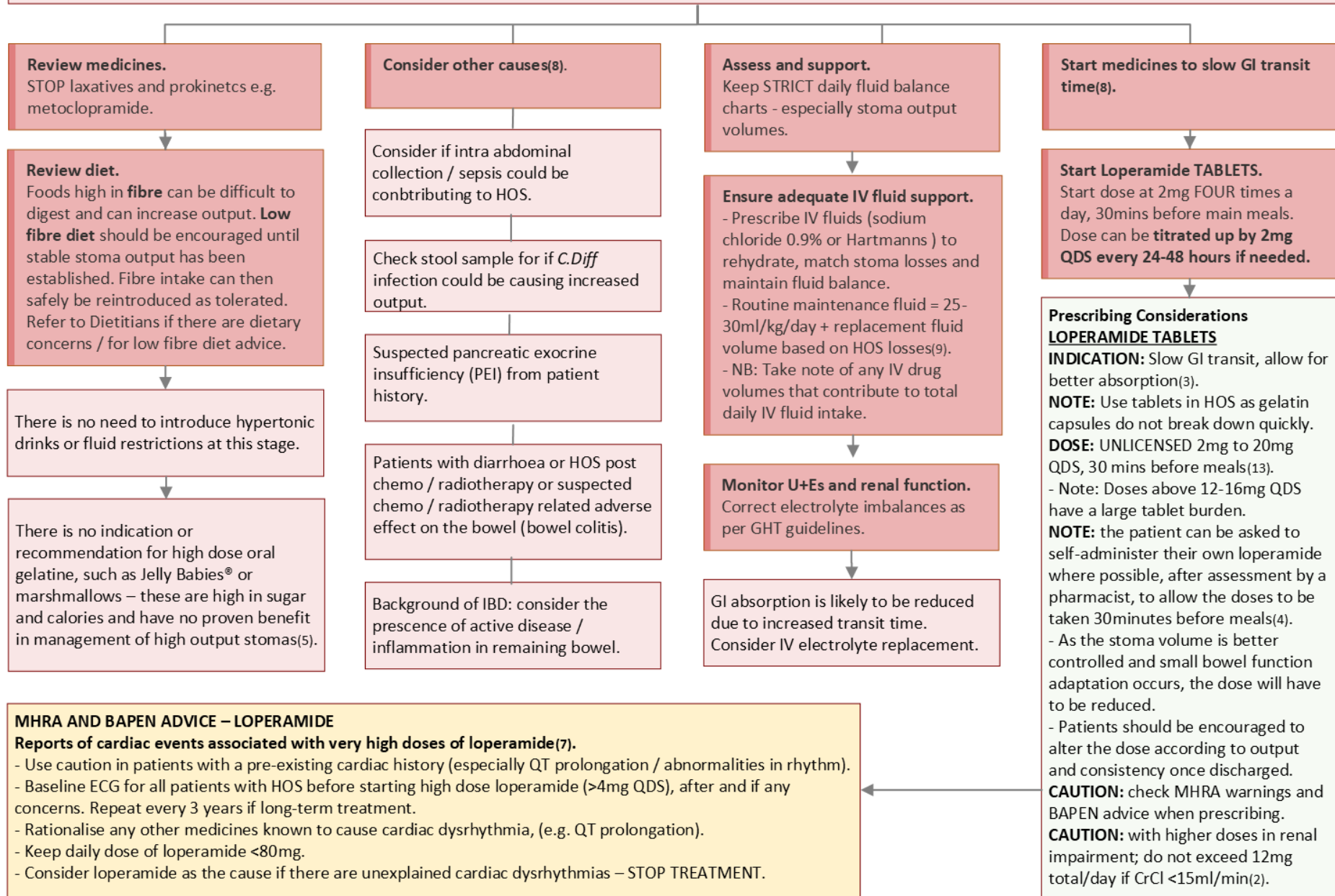
- **Stoma Care Specialists Nurses** – ensure there is a referral to stoma care nurses for stoma products review.

**Stoma Care Services:** <https://www.gloshospitals.nhs.uk/our-services/services-we-offer/stoma-care/>

# A

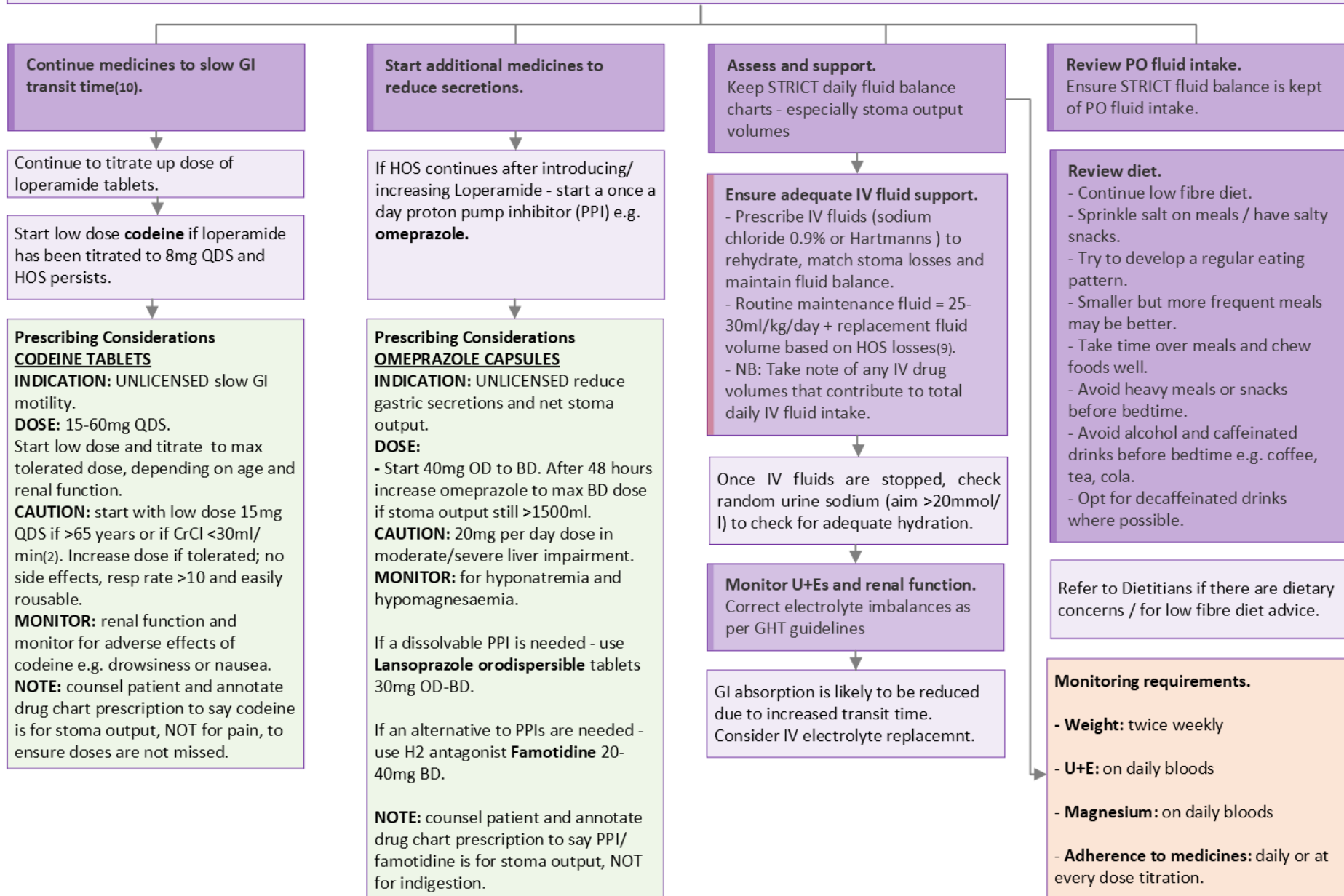
## INITIAL 48 hour management of HOS

**AIM:** stoma output approximately 1000ml **and** able to maintain own hydration, fluid and electrolyte balance with PO intake.



**B****CONTINUED management of HOS: patients who continue to have HOS despite escalating loperamide doses.**

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**Fluid restriction + St Mark's oral rehydration solution(10).**

- Oral rehydration solutions (ORS) have a high salt content (sodium  $\geq 90\text{mmol/L}$ ), e.g. St Mark's Solution and Dioralyte 'double strength'(1).

Introduce St Mark's ORS 1000ml per day and restrict PO intake of other fluids to 1000ml per day.

Review compliance with fluid restriction daily. Review effectiveness of fluid restriction after 48 hours.

If further PO fluid restriction needed:

- Max St Mark's fluid restriction to 1500ml per day.
- Min restriction of other PO fluids to 500ml per day.

If St Mark's ORS not tolerated due to palatability or errors in mixing – try **Dioralyte 'double strength'** (NOTE: see potassium content below).

**Prescribing consideration**

**ST MARKS ORAL SOLUTION(1)**

**INDICATION:** UNLICENSED ORS, 1000-1500ml per day with an oral restriction of other fluids.

**RECIPE:** Information for mixing St Mark's can be found at <https://www.gloshospitals.nhs.uk/your-visit/patient-information-leaflets/st-marks-formula-electrolyte-drink/>

**Prescribing consideration**

**DOUBLE STRENGTH DIORALYTE ORAL SOLUTION**

**INDICATION:** UNLICENSED dosing ORS. To drink 1000-1500ml per day as part of an oral restriction of other fluids.

**RECIPE:** 10 sachets dissolved in 1000ml of water.

**CAUTION:** when assessing suitability for patient, note that each 1000ml contains **40mmol potassium**.

**MONITOR:** U+Es and renal function.

**Consider starting octreotide to further reduce secretions if HOS persists(11).**

- Only start if initial management with first line anti-secretory/anti-motility medicines has been unsuccessful.
- Somatostatin analogues reduce output via a reduction in gastric, salivary, pancreatic and biliary secretions, while also delaying gastric emptying and slowing GI transit.
- Octreotide for HOS is unlicensed but can be a useful as a last line addition to the management of HOS if there is a large secretory output (PO fluid restriction is of minimal benefit)(12).

**OCTREOTIDE SUBCUTANEOUS INJECTION**

**INDICATION:** UNLICENSED - short acting somatostatin analogue to reduce net secretory output.

**DOSE:** Start at 50mcg s/c BD and titrate up to TDS after 24 hours. Can increase to 100mcg s/c TDS after a further 24 hours(6).

- Monitor for efficacy and increase dose by 50mcg TDS increments to max 200mcg s/c TDS, as tolerated.

**EFFICACY:** if no significant reduction in stoma output after 5-7 days at max tolerated dose, STOP octreotide.

Daily review by Surgical / Nutrition Support Team.

**CAUTION:**

- Monitor blood glucose BD-TDS whilst on octreotide as glucose regulation can be affected.
- Monitor heart rate - bradycardia has been reported; doses of any rate-controlling medicines may need to be adjusted. Monitor thyroid function and LFTs with prolonged use.
- STOP:** If severe abdominal pain / biliary colic develops. Acute pancreatitis and cholelithiasis / gall bladder events has been reported(6).

**NOTE:**

- Can cause stinging at injection site.
- Avoid abrupt withdrawal of treatment - wean dose to stop over 24-48 hours.
- Octreotide may affect absorption of dietary fats; faecal fat excretion may increase.

**Discharging the patient on a somatostatin analogue.**

- If somatostatin analogues are effective and required to continue on discharge, please discuss options with the Nutrition Support Team /ward pharmacist.
- Self administration of short acting octreotide pre-filled syringes or conversion to a long acting somatostatin analogue can be considered.
- Please note: a long acting preparation is not always possible as it requires patient specific local funding and communication with the GP.
- A change in formulation can destabilise current HOS control in the short term.

## References

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