Pancreatic surgery, enteral tube feeding & enzymes

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Introduction

- Statistics
- Anatomy
- Surgical treatment
- Enteral feeding
 - PERT
 - Evidence
 - What happens elsewhere

Pancreatic cancer



- 7 in 10 patients do not have any chemotherapy, radiotherapy or surgery
- 1 in 10 patients will go on to have curative surgery

Resectable pancreatic cancer





•Stage 1A means that the cancer is smaller than 2cm.

•Stage 1B means that the cancer is larger than 2cm – but is still contained in the pancreas.

•Stage 2A cancer is larger than 4cm and started to grow outside the pancreas, but has not spread to the lymph nodes.

•Stage 2B means the cancer has spread to nearby lymph nodes.

Anatomy



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Unresectable pancreatic cancer – palliative bypass









Gastrojejunostomy

Hepaticojejunostomy

Unresectable pancreatic cancer

Duodenal stent





Biliary stent



Anatomical areas



Whipples/Pylorus Preserving Pancreatico - Duodenectomy (PPPD)



PEI

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Tran, Lanschot, Bruno & van Eijeck. Pancreatology. 2009:9:729.

Distal Pancreatectomy





PEI incidence

16-60% pre-op 20-80% post-op

Varied results:

- Amount of pancreas removed/remaining
- Testing methods eg Feacal elastase, onset of steathorrea, prescription of enzymes
- Presence of symptoms malabsorb up to 55g before!
- Variable timescales pancreatic atrophy later on

Total Pancreatectomy





PEI incidence and malnutrition

Distal pancreatectomy	16-60% pre-op 20-80% post-op	Speicher & Traverso (2010) Phillips (2015)
Pancreacticoduodenectomy	22-45% pre-op 56-98% post op	Matsumoto & Traverso (2006) Phillips (2015)

- Loss of functional parenchyma
- Asynchrony of enzymes
- Oedema/obstruction at anastomosis
- Ph differentials

Loss of duodenum:

- Nutrient absorbtion
- Cholecystokinen secretion
- Bile flow
- Dumping
- Rapid transit

- <u>Effects of surgery</u>
 - Raised REE
 - Increased protein turnover
 - Reduced appetite
 - Pain, nausea, sickness
 - Drains
 - Medications
 - ?Infection

Nutrition support - enteral feeding

Enteral feeding - a key route of nutrition support in pancreatic disease:

Malignant disease -

- Prior to surgery
- Following surgery (eg. FTT)

Enteral feeding

How do we use PERT alongside enteral feeding?

- ESPEN (2006) recommend peptide feeds in pancreatic disease but patients still malabsorb
 - Therefore enzyme replacement therapy needed
 - PERT predominently designed for oral administration
 - ...little evidence to support/guide practice...
- Literature
 - International variability standards, practices, health insurance
 - Different EN formula
 - Different enzyme preparations, inc different doses
 - In-vitro studies
 - CF populations
 - Pomombor Coal: Pight time right place right pH

Evidence...

Ferrie (2011) (Austrailia) -

- Jejunal tubes: Open capsule, crush microspheres (remove coating), activate with Na bicarb 8.4%, flush,
- or, or dissolve uncrushed microspheres in Na bicarb for ~ 20-30mins, flush
- add directly to enteral feed

However:

- Crushing granules not advised in UK (Handbook of Drug Administration via Enteral Feeding Tubes, 3rd Ed. 2015)
- Time and labour intensive when patients require regular doses
- Unlicensed use
- Reduced enzyme effect with crushing and dissolving/activating
- <u>Gastric tubes</u>: Open capsule, maintain enteric coating, suspend in thickened acidic fluid eg. "nectar consistency fruit juice", administer

□ However:

- Consistency of fluid is key to avoid blocked tubes
- Tube lumen size: 10 & 12 Fr use low dose enzymes (eg. 5000IU not in UK, Hollander 2015 recommends no smaller than 16Fr using beads 0.71-1.6mm in Creon 24000u) = varied advice



aintain the



Figure 1. Effect of thickened fluid in preventing tube blockage. Behavior of enzyme microspheres when given in water (left) versus thickened fluid (right).



Locally

 Change from Creon, dissolved in Na bicarb, flushed 2-4hourly to Pancrex V mixed with water in gastric and jejunal tubes

□ Why?

- Cost
- Easier
- Less labour intensive
- Positive feedback from nursing staff!

Locally

Pancrex V	Lipase (BP units)
½ level 5ml teaspoon	25000
1 level 5ml teaspoon	50000
1&1/2 level teaspoons	75000
2 level 5ml teaspoons	100000

Directions:

- 1. Stop feed
- 2. Flush tube with water
- 3. Add prescribed dose of Pancrex V to a pot
- 4. Add 15mls water
- 5. Stir to disperse the Pancrex V powder
- 6. Draw into syringe and administer via feeding tube
- 7. Add further 15mls water to pot to ensure residual Pancrex V is dispersed
- 8. Draw into syringe and administer via feeding tube
- 9. Flush tube with water
- 10. Restart feed immediately

Starting doses:	NG	2tsp = 100 000u
	NJ	1½ tsp = 75 000u

Adding enzymes to feeds

- Used regularly in some centres in pancreatic malignant and benign disease
- Positive results anecdotally:
 - Less diarrhoea/improved frequency
 - No adverse effects
 - Feeds can split (esp Peptisorb)
 - Hanging times
 - Labour intensive requires good team understanding
 - Recorded results: improved wound healing, increased insulin reqs, less hypos, increased GS/weight (Phillips, Berry & Gettle 2018)

Future

- National online survey of using pancreatic enzymes alongside enteral feeds – watch out!
- Novel products/systems:
 - □ PERT cartridges not in UK
 - Relizorb (FDA approved) small plastic cartridge containing lipase, connects to EN giving set, hydrolyses fat as feed infuses
 - □ 1 cartridge per 500mls
 - Max 2 cartridges in 24 hours
 - Max rate 12omls/hr
 - Not compitable with feeds with soluble fibre
 - Poor results with TwoCal HN



(Phillips, Berry & Gettle 2018)

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Thank you!

Questions...

Evaluations...

Lunch!