

Small bowel tumours

اینه هاد اهم ارتی لازم نعرفه ایک الرکور اکور علیه آخراعها موری این هاد اهم ارتی لازم نعرفه

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- Only 1 to 2 per cent of malignant alimentary tumours.
- Equal between man and women
- 100 times less frequent than in the stomach, oesophagus, or colorectum
- Benign lesions are more common distal, while Adenocarcinoma is more common proximal.

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- Small intestinal tumors may originate in cells of the:
 - epithelium
 - adenomas,
 - adenocarcinomas or
 - carcinoids
 - lymphatic tissues
 - lymphomas
 - mesenchymal or neural elements
 - gastrointestinal stromal tumors
 - leiomyomas
 - lipomas
 - hemangiomas
 - neuromas
 - sarcomas

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Risk factors



- ► Familial adenomatous polyposis, → less common than colon CA
- Hereditary nonpolyposis colorectal cancer (HNPCC),
- Peutz-Jeghers syndrome,
- Crohn's disease,
- Gluten-sensitive enteropathy (celiac sprue),
- Biliary diversion (e.g., previous cholecystectomy).

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Presentation



- Sixth and seventh decades of life
- Benign tumours are found incidentally at laparotomy or autopsy
 - vague symptoms, absence of clinical signs, the difficulty in investigating much of the small bowel
 - nausea, dyspepsia, epigastric discomfort, fatigue, bloating and weight loss, to haemorrhage or obstruction
 - Haemorrhage: occult or major bleeding ...
 - palpable abdominal mass, perforation, fistula formation, intussusception or intraperitoneal haemorrhage

all non specific

CLINICA

٩L	PRESEN	IAIION	OF	PRIMARY	SMALL	ROMEL	TUMORS

Signs and symptoms	Frequency (%)				
BENIGN NEOPLASMS					
Asymptomatic	47–60				
Abdominal pain	24–50				
Acute gastrointestinal he	morrhage 29–44				
Anemia	28–58				
Intermittent obstruction	12–28				
* Asymptomatic	6–12				
*Abdominal pain	62–83				
*Weight loss	38–55				
Nausea/vomiting	23–64				
Acute gastrointestinal hemorrhage 6–31					
Anemia	12–38				
Abdominal mass	5–32				

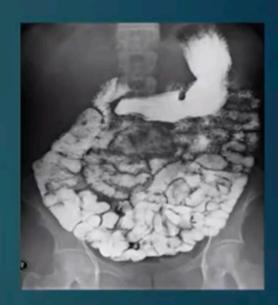






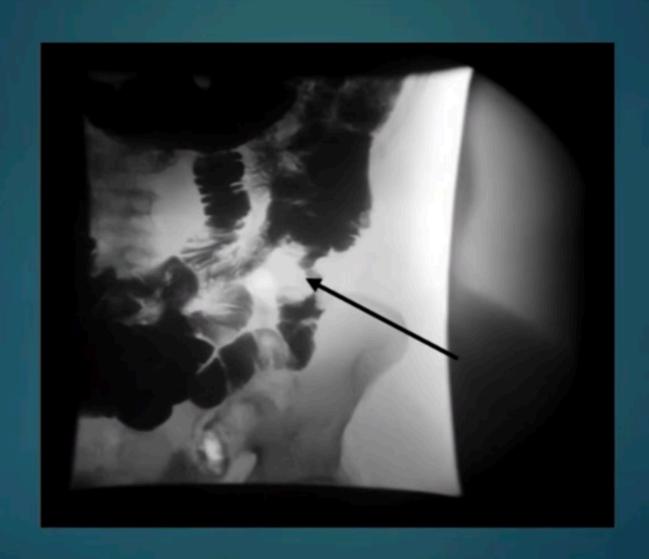
- Contrast Studies: small bowel follow through
- Endoscopy
- CT / MRI
- Angiography through the vessels
 Capsule endoscopy

Ssmall capsyle with a camera





small bowel follow through

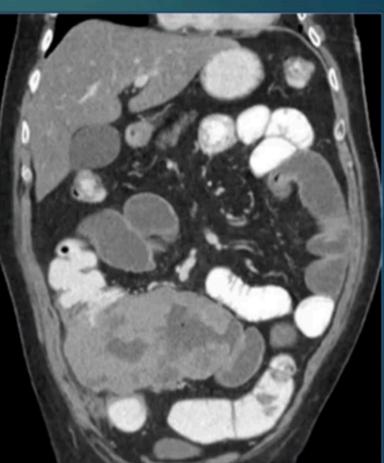


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CT entorography







thickening of the wall bowel

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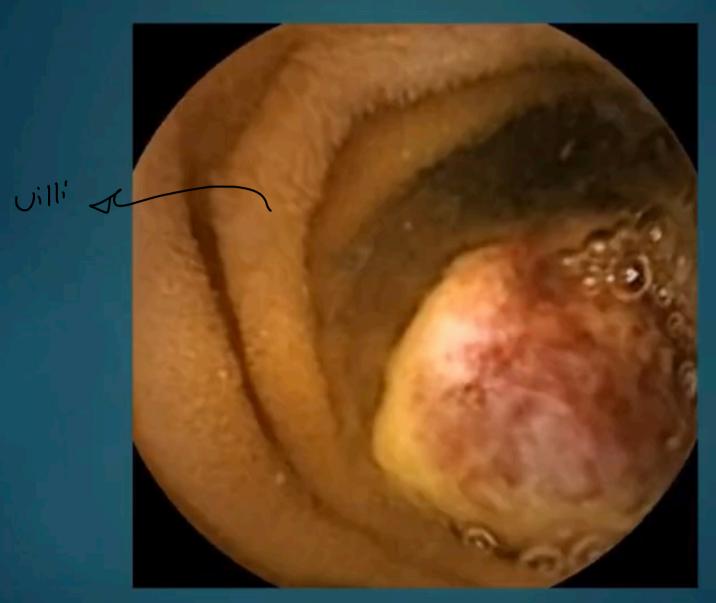


tumor in the right Iliac fossa

Angiogram

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Capsule endoscopy

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Pathological classification

- Not straightforward (not very clear)
 - Benign neoplasms
 - Malignant Tumours

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Adenoma - benign and originating from the epithelium

- True adenoma, Villous adenoma and <u>Brunner gland</u> adenoma.
- 20% in doudenum, 30 % in jejunum and 50% in ileum.
- Villous adenomas more common in Duodenum and less common in distal small bowel.
- Most common presentation is <u>asymptomatic</u> ,Abdominal pain, obstruction, and occult (or overt) haemorrhage Obstructive jaundice.
- Malignant change increases with the size, site, and increases with the histological type (
- number of lesions as well as with the histological type (tumours situated in the periampullary region are more prone to malignancy)



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lesion in the 2nd part of the chrodenum

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Familial Polyposis syndromes

- Familial polyposis coli
- Autosomal-dominant inheritance of the mutated APC gene
- Thousands of adenomas in the colorectum
- polyps occur within the small intestine in 24 to 93 per cent
- only 2 to 12 per cent develop duodenal cancer
- Treatment: excise polyps / biopsy / follow up

SB for all putients _ short-bowel syndrome

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Other Benign neoplasms

- Brunner's gland adenoma (doudenum)
- Lipoma
- Neurofibromas
- Fibroma
- Vascular tumours
- Leiomyoma

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Malignant Tumours of Small Intestine (more likely to produce symptoms)

- Malignant neoplasms almost always produce symptoms
- The most common :pain and weight loss
 - Obstruction in 15- 35% of patients (adhesions and infiltration)
 - Diarrhoea and excess mucus
 - GI bleeding , anaemia

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Carcinoid tumours

- Originate in enterochromaffin cells (argentaffin cells)
- ▶ 0.7 per 100 000
- These tumours may occur in the
 - foregut (including the duodenum),
 - midgut (including the jejunoileum),
 - the hind gut.
- Midgut carcinoids characteristically secrete large amounts of 5-hydroxytryptamine (5-HT; serotonin), whereas foregut carcinoids secrete small amounts of this peptide

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Carcinoid tumours cont.

- Most common in ileum (last two feet)
- Multi-centric in 30 40 % of cases
- yellow in colour and appear in a submucosal or serosal position
- *slow-growing tumours
- fifth decade, and both sexes are affected equally.
- Most common presentation is pain. Chance pain.
- Most patients will have metastasized to LN / Liver
- Doudenal carcinoid can cause ulceration, obstruction, and jaundice

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Carcinoid **tumours** malignant potential

- Metastasis is related to size of tumor,
 - ▶ Less than 1cm tumor: 20 30 % risk of mets to LN and liver
 - ▶ 1-2 cm tumor: 60-80% LN and 20% Liver
 - ▶ More than 2cm: 80% LN and 40-50% liver
- Lesion less then 1cm can be adequately treated with local excision.
- Small bowel obstruction, mesenteric fibrosis and ischaemia.

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Carcinoid syndrome.

- Carcinoid syndrome refers to vasomotor, gastrointestinal, and cardiac manifestations induced by systemic circulation of a variety of peptides elaborated by carcinoid tumor
 - Diarrhoea, flushing ,wheezes, abdominal cramps , cardiac (Rt heart failure)
 - Most likely liver metastases or large tumor bypass the liver
- Elevated urinary levels of 5-HIAA measured over 24 hours with high-performance liquid chromatography are highly specific but not sensitive

Carcinoid Tumours Cont.



- Treatment
- Local disease : resection including draining Lymph nodes
- Metastatic disease:
 - Tumour debulking, resection, cryotherapy, radiofrequency ablation, hepatic artery embolization, or chemoembolization.
- Systemic therapy
 - Somatostatin analogs Octeriotide, Long-acting octreotide, lanreotide. political
- Cytotoxic chemotherapy: ineffective.



Adenocarcinoma

- Adenocarcinoma accounts for about 35% of small bowel tumors
- More common in proximal small bowel than distal. and minimally in the
- median age at presentation of 60 years
- Presentation according to site:
 - Non-specific, vomiting, pain, jaundice, obstruction, perforation. lote presentation because of non-specific symptoms
- Jejunal and ileal tumours are best treated by segmental resection including the regional lymph nodes
- The overall 5-year survival rate for jejunoileal carcinomas is 20 to 30 per cent. If there is no nodal involvement at operation, survival is increased to 50 to 70 per cent
- Chemotherapy is of little help.

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Adenocarcinoma cont.

- Adenocarcinoma developing with crohn's disease is more common in lleum
- 20 years younger
- male preponderance of about 70 per cent
- prognosis is very poor

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Gastrointestinal lymphoma

- 1 to 4 per cent of all primary gastrointestinal cancers
- 50 to 55 per cent of tumours occur in the stomach, 30 to 32 per cent in the small bowel
- Present with; obstruction, bleeding, anorexia and weight loss.
- 5th and 6th decade
- Most common in ileum
- Increase incidence in patients with Coeliac disease / immunodeficiency states (e.g., AIDS).
 - Worsening diarrhea, pyrexia, and local obstructive symptoms.
- Treatment is usually medical unless surgical complication.



Gastrointestinal Stromal Tumours (GIST)

- Arise from Connective tissue cells
- Most common mesenchymal tumour of the GI Tract.
- Benign or malignant. Size increase risk of malignant potential.
- Usually stomach followed by small bowel (jejunum >ileum)
- 50-70 years of age.

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GIST cont.

- Lymphatic spread is not common (just resect the fumor)
- Metastasis to liver or peritoneum
- Prognosis
 - Worse than in stomach and oesphagus
 - Tumour size : less than 2 cm diameter > low risk
 - Mitotic rate: less than 5 HPF >> low risk



Treatment



- Surgery . Excision with negative margin
- Tyronise kinase inhibitor (imatinib) in advanced cases 50% tumours shrinkage.
- Radio-resistant



End of notes

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