GASTROINTESTINAL BLOCK PATHOLOGY LECTURE 2017/2018

MalignantTumors of Intestine

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OBJECTIVES:

Colon cancer:

- Describe the epidemiology of colon cancer.
- Compare the pathology (gross and microscopic features) and clinical features of right-sided colonic adenocarcinoma and left-sided colorectal adenocarcinoma.
- Describe the relationship between prognosis and the various stages of cancer of the colon and rectum as noted in the TNM (tumor-nodes-metastasis) classification and staging system.
- Describe the relationship between carcinoembryonic antigen (CEA) and recurrence following resection of the primary tumor.
- Mention the significant of carcinoid tumor and its features

TUMORS OF THE SMALL AND LARGE INTESTINES

Carcinoma Carcinoid tumor Lymphoma

• Adenocarcinoma of the colon is the most common malignancy of the GI tract and is a major cause of morbidity and mortality worldwide.

- The small intestine accounts for 75% of the overall length of the GI tract, is an uncommon site for benign and malignant tumors.
- Among malignant small intestinal tumors, adenocarcinomas and well-differentiated neuroendocrine (carcinoid) tumors have roughly equal incidence, followed by lymphomas and sarcomas.

MALIGNANT TUMORS OF LARGE INTESTINE

Adenocarcinoma

- > Adenocarcinoma of the colon is the most common malignancy of the GI tract and is a major cause of morbidity and mortality worldwide.
- Constitutes 98% of all cancers in the large intestine.
- ▶ incidence peaks at 60 to 70 years of age

MALIGNANT TUMORS OF LARGE INTESTINE Adenocarcinoma

Predisposing factors

- IBD, adenomas, polyposis syndrome.
- Diet appears to play an important role in the risk for 2. colon cancer:
 - Low fibre diet.
- Alcohol

- High fat content.
- -

Reduced intake of vit A, C & E.

It is theorized that reduced fiber content leads to decreased stool bulk and altered composition of the intestinal microbiota. This change may increase synthesis of potentially toxic oxidative by-products of bacterial metabolism, which would be expected to remain in contact with the colonic mucosa for longer periods of time as a result of reduced stool bulk. High fat intake also enhances hepatic synthesis of cholesterol and bile acids, which can be converted into carcinogens by intestinal bacteria.

• Several epidemiologic studies suggest that aspirin or other NSAIDs have a protective effect. This is consistent with studies showing that some NSAIDs cause polyp regression in FAP patients in whom the rectum was left in place after colectomy.

ADENOCARCINOMA OF LARGE INTESTINE

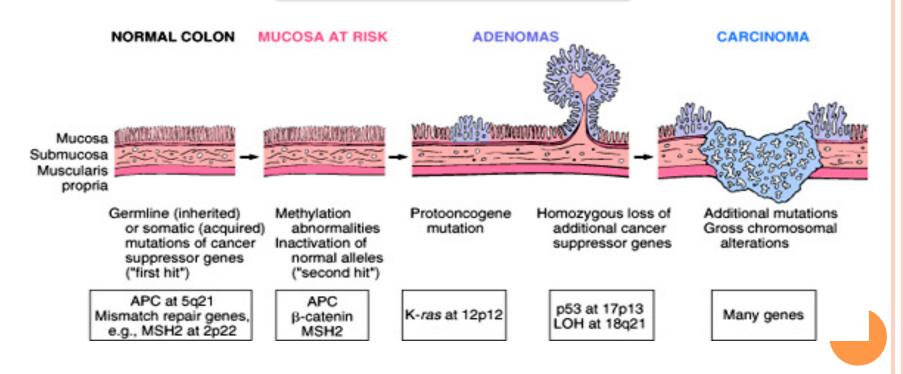
Carcinogenesis

- Two pathogenetically distinct pathways for the development of colon cancer, both seem to result from accumulation of multiple mutations:
 - 1- The APC/B-catenin pathway (85 %)
 - 2- The DNA mismatch repair genes pathway

ADENOCARCINOMA OF LARGE INTESTINE Carcinogenesis

- 1- The APC/B-catenin pathway (85 %)
 - chromosomal instability that results in stepwise accumulation of mutations in a series of oncogenes and tumor suppressor genes.

adenoma-carcinoma sequence



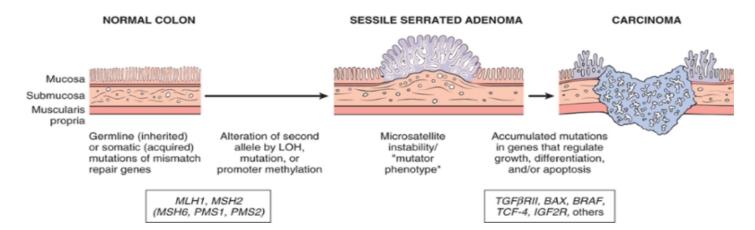
FAMILIAL ADENOMATOUS POLYPOSIS

• Hereditary mutation of the *APC* gene is the cause of familial adenomatous polyposis (FAP), where affected individuals carry an almost 100% risk of developing colon cancer by age 40 years.

MALIGNANT TUMORS OF LARGE INTESTINE ADENOCARCINOMA

- Carcinogenesis
- 2- The DNA mismatch repair genes pathway (These are referred to as MSI high, or MSI-H, tumors:)
- 10% to 15% of sporadic cases.
- There is accumulation of mutations
- Five DNA mismatch repair genes (MSH2, MSH6, MLH1, PMS1, AND PMS2)
- give rise to the <u>hereditary non polyposis colon</u> <u>carcinoma (HNPCC) syndrome</u>, known as Lynch syndrome.

HEREDITARY NON POLYPOSIS COLON CARCINOMA SYNDROME (DEFECTS IN MISMATCH REPAIR GENES_



HNPCC is caused by inherited germ line mutations in genes that encode proteins responsible for the detection, excision, and repair of errors that occur during DNA replication. This result in microsatellite instability and permit accumulation of mutations in numerous genes. If these mutations affect genes involved in cell survival and

proliferation, cancer may develop.

May produce abundant mucin that accumulates within the intestinal wall

COLORECTAL CARCINOMA

Morphology

- 70% are in the rectum, rectosigmoid and sigmoid colon.
- Left-sided carcinomas tend to be annular, encircling lesions with early symptoms of obstruction.
- <u>Right-sided carcinomas</u> tend to grow as polypoid, fungating masses, obstruction is uncommon.

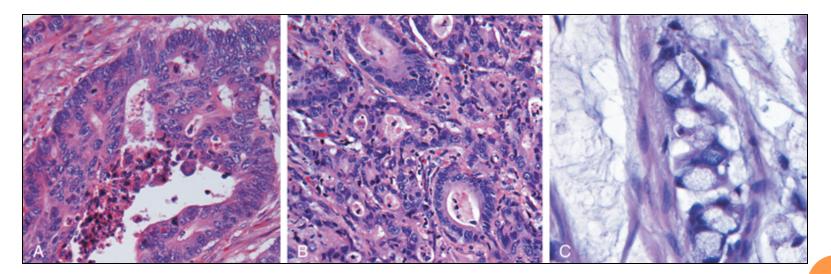




COLORECTAL CARCINOMA MORPHOLOGY

o Adenocarcinoma

• Mucinous adenocarcinoma secret abundant mucin that may dissect through cleavage planes in the wall.



SIGNS AND SYMPTOMS

- If located closer to the anus: change in bowel habit, feeling of incomplete defecation, PR bleeding
- A tumor that is large enough to fill the entire lumen of the bowel may cause bowel obstruction
- Right-sided lesions are more likely to bleed while left-sided tumors are usually detected later and could present with bowel obstruction.

Serum levels of carcinoembryonic antigen (CEA) are related to tumor size and extent of spread. They are helpful in monitoring for recurrence of tumor after resection.

TNM STAGING OF COLON CANCERS IS USED FOR STAGING

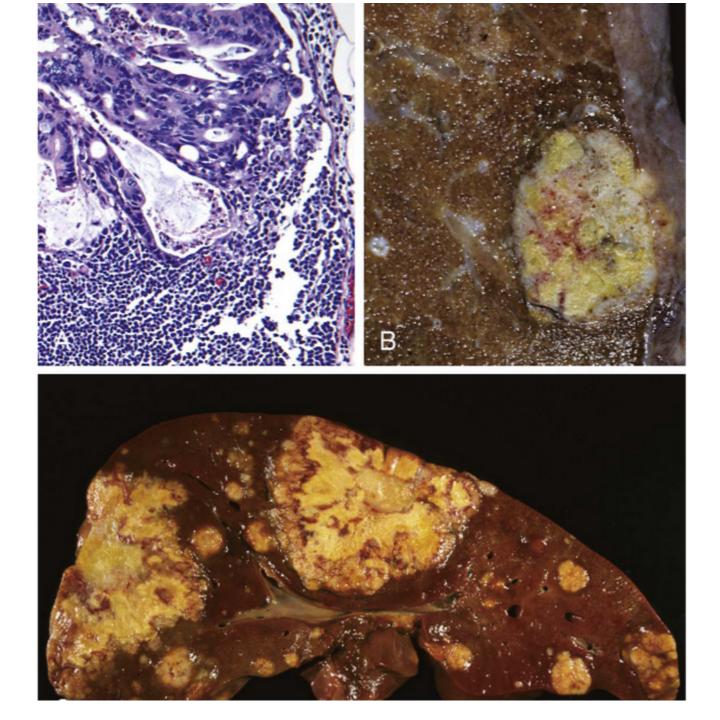
Table 14-8. AJCC Tumor-Node-Metastasis (TNM) Classification of Colorectal Carcinoma

Designation	Description				
Tumor					
Tis	In situ dysplasia or intramucosal carcinoma				
T1	Tumor invades submucosa				
T2	Tumor invades into, but not through, muscularis propria				
Т3	Tumor invades through muscularis propria				
T4	Tumor invades adjacent organs or visceral peritoneum				
Regional Lymph Nodes					
NX	Lymph nodes cannot be assessed				
N0	No regional lymph node metastasis				
N1	Metastasis in one to three regional lymph nodes				
N2	Metastasis in four or more regional lymph nodes				
Distant Metastasis					
MX	Distant metastasis cannot be assessed				
M0	No distant metastasis				
M1	Distant metastasis or seeding of abdominal organs				

AJCC, American Joint Committee on Cancer.

Table 14-9. AJCC Colorectal Cancer Staging and Survival

Stage*		Tumor-Node-Metastasis (TI	5-Year Survival (%)	
	Т	Ν	М	
l .	T1, T2	N0	M0	74
II				
IIA	T3	NO	M0	67
IIB	T4	N0	M0	59
Ш				
IIIA	T1, T2	N1	M0	73
IIIB	T3, T4	N1	M0	46
IIIC	Any T	N2	M0	28
IV	Any T	Any N	M1	6



COLORECTAL CARCINOMA

• The two most important prognostic factors are **depth of invasion** and the presence or absence of **lymph node metastases**.



MALIGNANT SMALL INTESTINAL NEOPLASMS

•In descending order of frequency:

- Carcinoid
- Adenocarcinomas
- Lymphomas
- Leiomyosarcomas.

SMALL INTESTINAL NEOPLASMS *Carcinoid Tumors*

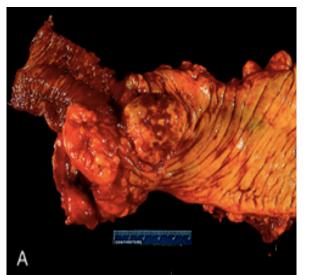
- Neoplasms arising from endocrine cells found along the length of GIT mucosa.
- The peak incidence: sixth decade, but they may appear at any age.
- They compose less than 2% of colorectal malignancies
- almost half of small intestinal malignant tumors:
 - 60 to 80% appendix and terminal ileum
- 10 to 20% rectum.

CARCINOID TUMORS BEHAVIOR

- Aggressive behavior correlates with:
- 1. Site of origin:
 - Appendiceal and rectal carcinoids infrequently metastasize, even though they may show extensive local spread
 - 90% of ileal, gastric, and colonic carcinoids that have penetrated halfway through the muscle wall have spread to lymph nodes and distant sites at the time of diagnosis, especially those larger than 2 cm in diameter.
- 2. Depth of local penetration
- 3. Size of the tumor

SMALL INTESTINAL NEOPLASMS Carcinoid Tumors Morphology

- A solid, yellow-tan appearance
- The cells are monotonously similar, having a scant, pink granular cytoplasm and a round-to-oval stippled nucleus.
- Ultrastructral features: neurosecretory electron dense bodies in the cytoplasm



SMALL INTESTINAL NEOPLASMS

Carcinoid Tumor

Clinical features

- Asymptomatic
- May cause obstruction, intussusception or bleeding.
- May elaborate hormones: Zollinger-Ellison, Cushing's carcinoid or other syndromes.

SMALL INTESTINAL NEOPLASMS

Carcinoid tumor

Carcinoid syndrome

- 1% of carcinoid tumor & in 20% of those of widespread metastasis
- Paroxymal flushing, episodes of asthma-like wheezing, right-sided heart failure, attacks of watery diarrhea, abdominal pain,
- The principal chemical mediator is serotonin
 The syndrome is classically associated with ileal carcinoids with hepatic metastases.

CLINICAL FINDINGS

• Due to serotonin and other bioactive compounds (e.g., histamine, bradykinin)

• Flushing of the skin (75%–90% of cases)

Due to vasodilation; may be triggered by emotion, alcohol, other foods

• Diarrhea (>70% of cases)

Increased bowel motility from serotonin

- Intermittent wheezing and dyspnea (25% of cases) Due to bronchospasm
- Facial telangiectasia
- Tricuspid regurgitation and pulmonary stenosis

Serotonin increases collagen production in the valves.

SEROTONIN AND DIARRHEA

• Patients with carcinoid syndrome often suffer from diarrhea, which has both a secretory and a motor component. The secretory component of carcinoid diarrhea is attributable to excessive serotonergic stimulation of submucosal secretomotor neurons; the motor component includes faster small bowel and colon transit and an exaggerated tonic response of the colon to ingestion of a meal

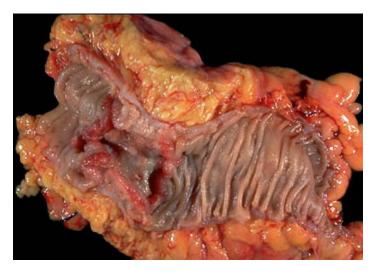
SMALL INTESTINAL NEOPLASMS

Lymphoma

- Most often low-grade lymphomas arising in mucosal-associated lymphoid tissue (MALT) lymphoma or high-grade non-Hodgkin's lymphomas of B cell type.
- May occur in any part of the intestine;
- The ileocecal region is a favored site for Burkitt's lymphoma.



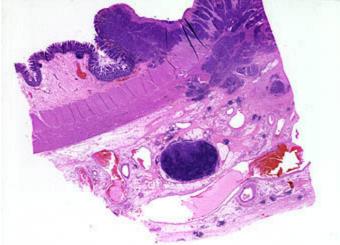




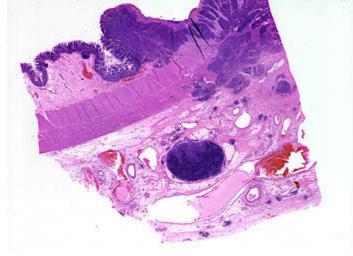
• Left colon, carcinoma - Gross, mucosal surface

This specimen from the left colon shows an annular, encircling, and constricting cancer. The margins of the cancer are heaped-up and firm, and the midregion is ulcerated.

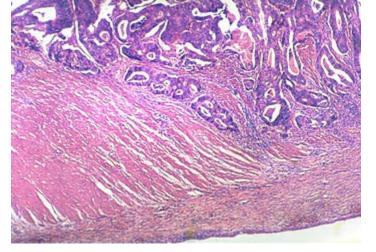
Left-sided colon cancers come to attention by producing occult bleeding and changes in bowel habits (i.e., constipation and cramping in the left lower quadrant).



- The carcinoma is composed of irregular glands infiltrating the muscularis propria, serosa, and mesentery. The TNM classification is based on the extent of invasion, number of lymph nodes involved, and extent of metastatic involvement. The deeper tumor extends into the muscularis propria, and as lymph nodes become involved, the prognosis worsens.
- T—extent of invasion
- N—number of lymph nodes involved
- M—extent of metastatic involvement



- Assuming this patient did not have lymph node metastasis, what stage is this carcinoma? The TNM stage for the current case would be T3N0MX.
- T3—extends through the muscularis propria
- N0—no lymph node involvement
- MX—extent of metastatic involvement unknown



• malignant glands of an adenocarcinoma of the colon infiltrating the muscularis propria.

What is the mode of spread of this cancer? Colonic carcinomas spread by local extension to adjacent structures. The favored sites of metastases are regional lymph nodes, liver, lungs, and bones.



Cecal adenocarcinoma

Tumors in the proximal colon tend to grow as polypoid, fungating, ulcerating masses. Obstruction is uncommon. About 25% of colon carcinomas are located in the cecum or ascending colon. Note the adjacent pedunculated adenomatous polyp. Most colon cancers develop from adenomatous polyps (the adenoma-carcinoma sequence).

Cecal and right colon cancers most often come to clinical attention by the appearance of fatigue, weakness, and iron-deficiency anemia.



DIVERTICULOSIS

