

## GI Surgical Oncology



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# Chapter 1

## Overview

### 1.1 Absences

Please notify Dr Hill, Dr Salo and Dr Squires before the beginning of the rotation if you will be away during the month. This includes vacations, meetings, interview trips, and other absences.

### 1.2 Communication

Please use Halo for messaging service attendings rather than SMS. Please check our status prior to messaging nights and weekends. If we are listed as unavailable, please contact another Surgical Oncology attending or the CMC “GI MIS Blue Surgery Attending Colorectal Onc” attending on call.

### 1.3 Inpatients

By in large, service attendings all wish to know about major changes in the status of our own patients. For most of the issues for which you would need to contact an attending at night, we would prefer that you HALO us directly rather than the on-call person. This desire is 24/7 (unless Halo says that attending is “off”).

### 1.4 ER admits

We would ask to use a combination of communication for ER admissions. All patients must be discussed with an attending. • If the patient is stable, does

not need surgery, etc. then we would ask you to contact the attending on call for GI MIS Blue Surgery Attending Colorectal Onc. Please contact them according to that attending's preferences (page, text, call, etc). • If the patient is unstable, may need surgery or will have ongoing and/or have intensive management needs the following day please HALO that patient's surgical oncology attending directly. The difference is that this patient is sick - we would like to know about all of our sick patients. If the Surg Onc attending is listed as "off" within Halo then please contact the GI MIS Blue Surgery Attending Colorectal Onc on-call attending.

## 1.5 Medical Records

Completing medical records in a timely fashion is critical for patient safety, billing, and compliance. Timeliness also demonstrates an understanding of how the world of surgery for which residents are being prepared functions.

## 1.6 Operative Logs

Completion of operative logs is critical for board certification of the individual resident but also has implications for the appropriate assignment of residents to surgical rotations AND impacts the ability of the residency to maintain accreditation and recruit resident candidates. Residents who find it difficult to find time to maintain operative logs may find themselves excused from the operating room to complete them. Residents are expected to complete operative logs within two weeks of the end of the rotation.

## 1.7 Case Assignment

The senior resident will be expected to make case assignments for junior residents and students. It is not necessary to split the month by attending - splitting by case is acceptable as well. We also expect that both residents know all the patients rather than just for one attending. This helps with nursing questions, etc.

## 1.8 Clinic

Clinic is an important part of a surgeon's education, where decisions are made regarding diagnostic workup, patient evaluation, and treatment planning. The expectation is that all residents on the service attend clinic once per week.



## **1.9 Work Hours**

If the service workload jeopardizes your ability to abide by the work hour restrictions, you must notify an attending so that arrangements can be made. The service attendings are committed to abiding by work hour restrictions.



## Chapter 2

# CMC Inpatient

Colorectal Surgery (Davis/Kasten) and GI Surgical Oncology (Hill/Salo/Squires) will cover Pineville and CMC. For efficiency, the services at each hospital will merge for patient care. Each patient will continue to have an attending surgeon, but rounding and inpatient care will be provided by the service.

In general Dr Kasten will round on his own patients at CMC.

### 2.1 Admissions

Admitting Provider: "CMC GI SURGICAL ONCOLOGY"

List Attending Surgeon in addition

Patient List is CMC GI Surgical Oncology

### 2.2 Rounds

Attending rounds for both services (CR and SurgOnc) start at 6am in STICU or 11T. Dr Salo rounds M-Tu and Drs Squires and Hill round W/Th/F alternate weeks

### 2.3 Resident Epic teams:

- CMC LCI GI SURGICAL ONCOLOGY NEW CONSULT
- CMC LCI GI SURGICAL ONCOLOGY CALL
- CMC LCI GI SURGICAL ONCOLOGY TEAM

- CMC AH COLORECTAL SURGERY CALL
- CMC AH COLORECTAL SURGERY CALL - DAVIS/KASTEN

Residents will be assigned to Epic teams by schedule. It is critical that you notify service attendings before the start of the month to adjust the resident call schedule. Each “shift” is 5:50am to 6pm. At 6pm the resident Halo Teams will be forwarded to the night team.

Please append a text block to the bottom of each progress note specifying the Halo Team for that patient to facilitate communication from nursing.

## 2.4 Attending Epic teams:

- CMC AH COLORECTAL SURGERY ATTENDING
- CMC LCI GI SURGICAL ONCOLOGY ATTENDING

Monday through Thursday (24hr), please contact the attending surgeon for each patient. For Surgical Oncology, please use Haiku. For Colorectal, please use phone.

Friday and Weekend: Epic Secure Chat “CMC GI MIS Blue Surgery Attending Colorectal Onc” for new admissions. Please keep Surgical Oncology attending informed of inpatient issues.

## 2.5 Consults

Established patients and directed should be discussed with the attending surgeon.

Unassigned Colorectal: “CMC AH COLORECTAL SURGERY ATTENDING”

Unassigned Surgical Oncology: “CMC LCI GI Surgical Oncology Attending”

In general, benign colorectal consults are staffed by Dr Davis. Colorectal malignancies are staffed as below. Esophageal and GE junction staffed by Dr Salo. Adenocarcinoma of distal stomach: Drs Salo/Squires. Gastric GIST: Drs Hill/Salo/Squires. Squires/Hill alternate week

	Mon	Tues	Weds	Thu	Fri
CR Malig	JSH	MHS	MHSJSH	JCS	MHSJSH
GI Surg Onc	JSH	MHS	MHSJSH	JCS	MHSJSH

## 2.6 Postop Clinic Appts

Postoperative patients are generally seen for a Transition of Care visit within the first week

Discharge appointments are made by sending a message in Canopy the evening prior (preferred) OR the morning of discharge before 8am to:

- LCI CMC GI, Clerical
- Marsha Sukhdeo (Salo and Squires)
- Sara Gaddy (Salo and Squires)
- Rebecca Wicks (Hill)
- Brenda Prieto
- Brandon Galloway

Please include the following information in the Canopy Message:

- Name of attending
- Ward from which the patient is being discharged
- Desired date for appointment
- Need for Wound Ostomy RN appointment at same time (essential for new stomas)
- Need for bloodwork at first
- Other studies to be done after discharge
  - Upper GI
  - Chest X-ray
  - Modified Barium Swallow

If messages cannot be sent in time, clinic schedulers can be reached at: Marsha Sukhdeo (Salo and Squires): 980-442-6110 or Carmen Davis (Hill): 980-442-6183.

If schedulers are not available, clinic RNs can be reached at: (Hill) 980-442-6146 or (Salo and Squires) 980-442-6143.

For patients likely to go home over the weekend or holidays, please plan to send a canopy message before 3pm on Friday or the day prior.

The scheduler will respond with a message to the discharging resident AND to the ward CNL with the appointment time, which can be included within the discharge summary. Copies of the message will also be sent to clinical nurse leaders:

- 11Tower: Sharon Hood
- 6Tower: Amy Peterson

## 2.7 Conferences

- GI Tumor Planning Conference Monday 7-8am
- Resident Teaching Conference Tuesday 7-8am 5th floor LCI II. Please review the upcoming clinic schedule and choose a case to present.
- Bone and Soft Tissue Conference Friday 7-8am

## Chapter 3

# Pineville Inpatient

### 3.1 Rounds

- Monday - Dr Squires
- Tuesday - Dr Hill

Starting time for rounds is variable from day to day.

Maddie Georgino will help organize work and timing of rounds, etc.

### 3.2 Resident Halo teams:

Pine AH Colorectal Surgery 1st Call Pine LCI GI Surgical Oncology 1st Call

Residents will be assigned to Halo teams by schedule. It is critical that you notify service attendings before the start of the month to adjust the resident Halo schedule. Each “shift” is 5:50am to 6pm. At 6pm the resident Halo Teams will be forwarded to the night team.

Please append a text block to the bottom of each progress note specifying the Halo Team for that patient to facilitate communication from nursing.

### 3.3 Attending Halo teams:

Pine AH Colorectal Surgery Attending Pine LCI GI Surgical Oncology Attending

Monday through Thursday (24hr), please contact the attending surgeon for each patient. For Surgical Oncology, please use Halo. For Colorectal, please use phone.

Friday and Weekend: Halo “CMC GI MIS Blue Surgery Attending Colorectal Onc” for new admissions. Please keep Surgical Oncology attending informed of inpatient issues.

### 3.4 Consults

Established patients and directed should be discussed with the attending surgeon.

Unassigned Colorectal: “Pine AH Colorectal Surgery Attending”

Unassigned Surgical Oncology: “Pine LCI GI Surgical Oncology Attending”

### 3.5 Postop Clinic Appts

Postoperative patients are generally seen for a Transition of Care visit within the first week

Discharge appointments are made by sending a message in Canopy the evening prior (preferred) OR the morning of discharge before 8am to:

- Hale Mock
- Syreeta McNair
- Madeline Georgino

Please include the following information in the Canopy Message:

- Name of attending
- Ward from which the patient is being discharged
- Desired date for appointment
- Need for Wound Ostomy RN appointment at same time (essential for new stomas)
- Need for bloodwork at first visit
- Other studies to be done after discharge
  - Upper GI
  - Chest X-ray
  - Modified Barium Swallow

For patients likely to go home over the weekend or holidays, please plan to send a canopy message before 3pm on Friday or the day prior.



## **3.6 Conferences**

- GI Tumor Planning Conference Monday 7-8am
- Resident Teaching Conference Tuesday 7-8am 5th floor LCI II. Please review the upcoming clinic schedule and choose a case to present.
- Bone and Soft Tissue Conference Friday 7-8am



## Chapter 4

# Rounds

Please plan to update the attending of record about their patients in the morning.

The following format will help speed communication on rounds.

**ID:** One line description: “Mr Glenn: PostOp day 3 after low anterior resection”

**24 hour events:** Summary of important events in prior 24 hrs

**Systems-oriented Presentation:**

Neuro: Pain control, level of alertness, psychotropic meds, sedatives, and pain meds.

CardioVascular: Vital signs (normal OR cite the range of systolic blood pressures and range of heart rate). Heart rhythm. Cardiac meds. Most recent recommendations of cardiology consult.

Respiratory: Pulmonary exam, oxygen saturation, supplied oxygen, ventilator setting. Results of CXR.

GI: Diet, bowel function, NG output. Drain outputs can often be summarized unless they are unusually high or low (and ready to be removed. New finding of bile in any abdominal drain needs special emphasis. GI meds (eg protonix, Entereg). Tube feed formula, rate and duration (continuous or nocturnal). Status of C Diff tests. Results of JP drain amylase levels (gastroesophageal patients). Results of JP triglycerides or creatinine, if sent,

Renal: Urine output in 24 hours AND in most recent 8 hour shift. Presence (or absence) of Foley catheter and plans for removal, if present. Most recent creatinine. If diuretics administered, dosage and amount of urine output during the shift when it was administered. Most recent potassium in any patient receiving (or about to receive) furosemide (Lasix). Results of Mg and Phos if abnormal.

Heme: Hemoglobin, platelets, DVT prophylaxis. PLEASE CHECK THE MAR SUMMARY DAILY to be certain that the ordered DVT prophylaxis has been given.

ID: WBC, Tmax in past 24 hours, culture results.

Endo: Diabetic regimen, blood sugar range, and amount of sliding scale insulin administered in the prior 24 hours.

**Problem-Oriented Plan:**

Each of the patients problems are addressed with an assessment and plan. Pre-existing medical problems and postoperative complications need to be addressed in the plan

## Chapter 5

# Progress Notes

Progress notes should include a comprehensive summary of pertinent information for the patient's medical care.

A running summary of events is recorded in the Shared Hospital Course under Inpatient Workflow

Each day, this section is carried from note to note so that each progress note contains the cancer history and the daily events.

The Progress Note then includes the date and a one-line summary of the operation performed and any intraoperative complications or events which impact the post-operative care.

Each day, an additional line is added to the Shared Hospital Course which summarizes events for that day. This makes it possible to see within each Progress Note the pertinent events for the hospitalization. These events would include extubation, re-intubation, positive cultures, dates lines are inserted or removed, dates of removal of NG tubes and drains, and transfer to ward or re-admission to ICU. This chronology assists in treatment decisions ("how old is the IJ line" or "when did we start antibiotics?" or "when is the planned antibiotic stop date"?) but also makes the discharge summary much easier to prepare.

Assessment/Plan The medical problems currently being managed are addressed in the assessment.

### **Esophagectomy Events to be Documented:**

- Extubation date/time
- NG Removal date
- Chest tube removal date
- MBS date(s) and results (aspiration | penetration)

- ICU DC orders written
- ICU discharge (transfer to ward)

### Esophagectomy Complications to be Documented:

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N	Delirium
	Stroke
CV	New arrhythmia req Rx
	MI
R	Pneumonia (3 of fever   WBC   infiltrate   abx   sputum cx)
	Effusion req drainage
	Reintubation
	Atelectasis req bronchoscopy
	ARDS
	PE
	Ventilation >48 hours after leaving OR
GI	Anastomotic leak (medical rx   stent   surgery)
	Delayed gastric emptying req botox or NG >7d
	C Diff
GU	Urinary Retention
	Discharge with foley catheter
H	DVT req treatment
	Return to OR
	Return to ICU

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### Communication

Please add an addendum at the BOTTOM of each progress note which includes a means for contacting the team:

Please message “GI Surgical Oncology LCI CMC” via Haiku 24/7. Messages are automatically forwarded to the General Surgery Resident on Call evenings and weekends.

#Signout

### Evening Signout

The Handoff Tool should be completed for all inpatients, and the responsible attending designated. This tool is critical for the safe care of patients by the night team. If there are studies which are pending at the time of signout (CT scan, follow-up Hb), it is critical that a plan be in place for whom to notify with an abnormal or critical study. In general, Drs Hill and Salo are always available until 10pm. Attending notification plans (service attending vs covering attending) for unstable patients should be negotiated before nightfall.

### Weekend Signout

The chief resident is responsible for making certain that the weekend rounding resident is familiar with the patients, their problems, and the plan of care. A signout email should be prepared Friday afternoon and forwarded to the service attendings by 6pm for their review. This signout can then be edited with the attendings' notes and forwarded to the weekend rounding attending.





## Chapter 6

# Discharges

### Discharge Prescriptions

Prescriptions should be ideally be prepared the day prior to anticipated discharge and sent to the patient's pharmacy. According to North Carolina STOP guidelines, opioid prescriptions for postoperative patients should be for no more than a 7 day supply.

### Additional Appointments

If followup appointments in addition to surgical followup are needed, these should be designated on the discharge orders. Particularly:

- Primary Care Physician
- Cardiologist (if new cardiac medicines)
- Co-surgeons (Urology, Thoracic Surgery, GYN)

### Discharge Summary

The discharge summary documents important events and complications in the postoperative course and serves to inform the referring physician and primary physician about these events, but also serves as a blueprint for post-discharge treatment planning. Please recognize that the first post-operative visit may be with a resident who may be meeting the patient for the first time. Key items to include:

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N	Followup plan for chronic pain management Stroke
CV	Postop Arrhythmia?   MI?   CHF? If new cardiac meds: Who is managing medications If afib: CHADS score and anticoagulation plan
R	Pneumonia?   ARDS?   TRACH?

---

	Need for home oxygen?
	CXR needed at first postop visit?
GI	Delayed gastric emptying?   leak?   ileus?
	Tube feed regimen
	Diet at discharge (Low residue   Full liquids   Meds with thickened water   NPO)
	New stoma (ileostomy   colostomy)
	Wound care needs (VAc   Prevena)
GU	Urinary Retention
	Discharge with foley catheter
H	Complications: DVT   PE
	Anticoagulation Plan
Endo	Insulin regimen at DC (dose will be in med rec)
ID	Antibiotics at DC
	Return to ICU

---

### Communication

It is essential that discharge summaries be sent to the patient's primary MD and referring physician. Please review the initial consultation note for the names of providers involved in a patient's care.

## Chapter 7

# Education

### 7.0.1 Student EMR Notes Policy

Approved: Medical Education Division of Canopy Team 11/17/13

Students:

Should write Notes in the EMR as a part of their training. Students must sign and forward their notes. It is imperative that student notes not be saved and forwarded prior to signing the note or the authorship of the note will change. Signed notes should be forwarded to the supervising physician as soon as possible.

Supervising physicians:

Attending must sign the necessary documents per the medical staff bylaws (e.g., H & P's and Discharge Summaries). Attendings, residents, or fellows can cosign progress notes. Each attending should clarify who is responsible for signing progress notes.

Supervising physicians have 2 options:

Correct the note as needed, append an attestation, and sign within 24 hours. When appending the student's note, the supervising physician may utilize only the student's past medical history, family history, social history, and review of systems. This is the preferred model and the most efficient. Please see example below.\*

Correct the note as needed, attest with reference to supervising physician's own note, and sign within 24 hours.

Example Attending Attestation\*

I have discussed the presentation of the patient with the student. I have also independently seen and examined the patient. Please see my documentation below:

History: [As obtained by the provider]

Past History, Family History, Social History, and ROS: Please see the student's note above.

Physical Exam:

General: Well-developed, well-nourished

HEENT: Pupils equal react to light extra ocular motion intact, conjunctiva pink, anicteric sclera, oropharynx clear, moist mucosa

Neck: Supple, no meningeal signs

Lungs: Clear to auscultation bilaterally, bilateral breath sounds, normal effort

Cardiac: Regular rate normal rhythm, no murmur gallop or rub

Abdomen: Soft, benign, non-distended, non-tender, no mass

Extremities: Moves all 4 extremities, pulses 2+ and equal

Neurologic: Awake, alert, and oriented x3, cranial nerves II through XII are intact, moves all 4 extremities equally, speech is normal, affect pleasant

Skin: No rash

Labs, X-rays, Tests: [Pertinent comments regarding positives/negatives and interpretations]

Medical Decision Making: [MDM also should be attending documentation]

Electronically Signed: Dr. Supervision

## Chapter 8

# Clinic

Clinic is an essential part of the educational experience as this is where preoperative evaluation and surgical planning is done.

### 8.1 Salo Clinic

Notes are generated using the .LCICONCONSULT SmartPhrase template. The will automatically pull in the names of the cancer care team (medical oncologist, gastroenterologist). History from prior notes can be pulled in and placed in the Subjective section. The problems pertinent to the visit are designated.

As always, notes which are copied forward need to be carefully proofread to avoid errors. For instance, when a patient returns for a postop visit, it is important not to carry forward text from the preoperative note that states 'surgery next week.

It is important that authorship be clearly communicated, particularly with regard to assessment and plan. In many cases, the assessment and plan area straightforward and can be copied forward from the prior Assessment/Plan. In some cases, clinical decision-making involves a judgement call. This is incicated in Dr Salo's notes as 'I would recommend...' In these cases, please do no copy the assessment notes verbatim to avoid confusion about whose opinions are being quoted.



## Chapter 9

# Postop Care after Esophagectomy

### 9.1 Weaning Tube Feeds in Diabetics

As outpatients begin eating more orally, their tube feeds are reduced.

Weaning from 5 cans to 4 cans: Easiest method is to maintain the same schedule (16 hours) and reduce insulin dosage by 20%. For instance, the above patient who is on 5 cans at 75mL/hour x 16 hours is receiving 16N + 8R at start of tube feeds and 6 hours later. This patient could be weaned by reducing rate from 75mL/hour x 16 hours to 60mL/hour x 16 hours and reducing insulin to 12N + 6R at the start of tube feeds AND another dose of 12N + 6R after 6 hours.

Weaning from 4 cans to 3 cans: One option is to decrease the duration of tube feeds from 16 hours to 12 hours, while maintaining rate of 60mL/hour. In this case, the insulin dosage could be kept the same at the start of tube feeds, BUT the dose 6 hours after the start of tube feeds could be omitted.

Once patients are on 3 cans per night, further weaning can be accomplished by skipping tube feeds (and insulin) every other night in a “tube feed holiday”. This allows an evening of interrupted sleep and can tend to increase the appetite the morning after tube feeds are held.





# Postoperative Care



# Chapter 10

## Colectomy

### Clinic

- Opioid Cessation
- Smoking Cessation
- EtOH/Drugs of Abuse – Social Work
- Nutritional Evaluation
  - All Patients – Ensure for 3d preop
  - Poor nutrition – ?Delay surgery
- Preop Anesthesia/ERAS Class
- Expectations of Surgery:
  - Length of Stay 1-3 days
  - Diet (self-limiting)
  - Pain control (low-opioid)
  - Activity (OOB at 6am, OOB 3x/day)
- Bowel Prep – Abx and mechanical

### Preop Holding

- Colon PowerPlan (Hill)
- Antibiotic PowerPlan
- No PCN only for severe allergy
- Entereg (if no preop opioids)
- VTE prophylaxis
- Carbohydrate load (2hr preop)

### OR

- Goal-directed fluid administration
- 2L total in OR
- 3L total/first 24hrs
- Open procedures: No epidural

### Postop Day 0

- Goal-directed fluid administration
- OOB (Dangling not compliant)
- Diet
  - Low Residue diet
  - Ensure Supplements
- Teaching
- Gum, Mag & Entereg
- Pain Management
  - PCA
  - Tylenol 1gm q6 ATC
  - Gabapentin 300mg TID
  - Tramadol PRN
  - Resume all baseline pain meds
- Home Medications
  - Resume all home medications
  - No therapeutic anti-coagiation
  - Diabetes medicines
    - \* Prefer to resume all oral diabetes meds
    - \* Sliding-scale insulin ordered for all diabetics

### Postop Day 1

- Labs: K+ and CBC only
- d/c “pre” plan & colon visit
- Heparin lock IV
- Remove Foley
- d/c PCA (unless laparotomy)
- Out of bed > 6 hrs
- Diet: low residue
- Pain management
  - 1) d/c PCA
  - 2) Tylenol
  - 3) Gabapentin 300 tid
  - 4) Tramadol

- 5) Home meds
- 6) Oxy if lots of pain
- Discharge planning
- 1) Possible home today!(25% will go home POD1)
- 2) Check patient 2-3PM

- Afternoon rounds
- 1) Possible home today!
- 2) Check patient 2-4PM
- 3) Ambulation 2x's by PM
- 4) Patient education

## **Postop Day 2**

- No IV fluids unless indicated
- No labs unless indicated
- OOB >6 hrs
- No PT unless going to rehab or SNF (ask Hill first)
- Pain management
  - Same as POD#1
- Discharge planning
  - Consider early D/C (median LOS=2d)
- Otherwise plan
  - Patient
  - Nursing
- Afternoon rounds
- 1) Possible home today!
- 2) Check patient 2-3PM
- 3) Ambulation 2x's by PM
- 4) Check for dehydration
- 5) Patient education
- 6) Give estimated date of discharge

## **Postop Day 3-5**

- IVF

- Consider bolus of IV fluids
  - Consider maintenance IV if we feel won't resolve soon
- Labs
  - No labs unless indicated
  - Consider ordering QOD Chem7 if prolonged ileus
  - OOB >6 hrs
  - No PT unless likely to go to rehab or SNIF (ask Hill first)
- Pain management
  - Same as POD#1
- Afternoon rounds
  - 1) Possible home today!
  - 2) Check patient 2-3PM
  - 3) Ambulation 2x's by PM
  - 4) Check for dehydration
  - 5) Patient education
  - 6) Give estimated date of discharge

# Chapter 11

## LAR + Ileostomy

### Clinic

- Opioid Cessation
- Smoking Cessation
- EtOH/Drugs of Abuse – Social Work
- Nutritional Evaluation
  - All Patients – Ensure for 3d preop
  - Poor nutrition – ?Delay surgery
- *Arrange Wound Ostomy Nursing*
- *Pre-approval for Home Health*
- Preop Anesthesia/ERAS Class
- Expectations of Surgery:
  - Length of Stay 1-3 days
  - Diet (self-limiting)
  - Pain control (low-opioid)
  - Activity (OOB at 6am, OOB 3x/day)
- Bowel Prep – Abx and mechanical

### Preop Holding

- Colon PowerPlan (Hill)
- *CCM order*
- *WOCN order*
- Antibiotic PowerPlan
- No PCN only for severe allergy
- Entereg (if no preop opioids)
- VTE prophylaxis

- Carbohydrate load (2hr preop)
- Confirm stoma marking
- Confirm CCM aware of ostomy

**OR**

- Goal-directed fluid administration
  - 2L total in OR
  - 3L total/first 24hrs
- Open procedures: No epidural

**Postop Day 0**

- Goal-directed fluid administration
- OOB (Dangling not compliant)
- Diet
  - Low Residue diet
  - Ensure Supplements
- Teaching
- Gum, Mag & Entereg
- Pain Management
  - PCA
  - Tylenol 1gm q6 ATC
  - Gabapentin 300mg TID
  - Oxycodone once at night
  - Tramadol PRN
  - Resume all baseline pain meds
- Home Medications
  - Resume all home medications
  - No therapeutic anti-coagiation
  - Diabetes medicines
    - \* Prefer to resume all
    - \* Sliding-scale insulin if needed
- *Wound Ostomy teaching*
- *CCM for Home Health for stoma care*

**Postop Day 1**

- Labs: K+ and CBC only
- d/c “pre” plan & colon visit



- Heparin lock IV
- Remove Foley
- d/c PCA (unless laparotomy)
- Out of bed > 6 hrs
- Diet: low residue
- Pain management
  - 1) d/c PCA
  - 2) Tylenol
  - 3) Gabapentin 300 tid
  - 4) Tramadol
  - 5) Home meds
  - 6) Oxy if lots of pain
- Discharge planning
  - 1) Possible home today!(25% will go home POD1)
  - 2) Check patient 2-3PM
- Afternoon rounds
  - 1) Possible home today!
  - 2) Check patient 2-4PM
  - 3) Ambulation 2x's by PM
  - 4) Patient education

## Postop Day 2

- No IV fluids unless indicated
- No labs unless indicated
- OOB >6 hrs
- No PT unless going to rehab or SNF (ask Hill first)
- Pain management
  - Same as POD#1
- *Wound Ostomy Teaching*
- Discharge planning
  - Consider early D/C (median LOS=2d)
- Otherwise plan
  - Patient
  - Nursing
  - *CCM for home health for stoma care*
- Afternoon rounds

- 1) Possible home today!
- 2) Check patient 2-3PM
- 3) Ambulation 2x's by PM
- 4) Check for dehydration
- 5) Patient education
- 6) Give estimated date of discharge

**Postop Day 3-5** - IVF - Consider bolus of IV fluids - Consider maintenance IV if we feel won't resolve soon - Labs - No labs unless indicated - Consider ordering QOD Chem7 if prolonged ileus - OOB >6 hrs - No PT unless likely to go to rehab or SNIF (ask Hill first) - Pain management - Same as POD#1 - Afternoon rounds 1) Possible home today! 2) Check patient 2-3PM 3) Ambulation 2x's by PM 4) Check for dehydration 5) Patient education 6) Give estimated date of discharge

## Chapter 12

# Abdominoperineal Resection

### Clinic

- Opioid Cessation
- Smoking Cessation
- EtOH/Drugs of Abuse – Social Work
- Nutritional Evaluation
  - All Patients – Ensure for 3d preop
  - Poor nutrition – ?Delay surgery
- *Arrange Wound Ostomy Nursing*
- *Pre-approval for Home Health*
- Preop Anesthesia/ERAS Class
- Expectations of Surgery:
  - Length of Stay 1-3 days
  - Diet (self-limiting)
  - Pain control (low-opioid)
  - Activity (OOB at 6am, OOB 3x/day)
- Bowel Prep – Abx and mechanical

### Preop Holding

- Colon PowerPlan (Hill)
- *CCM order*
- *WOCN order*
- Antibiotic PowerPlan

- No PCN only for severe allergy
- Entereg (if no preop opioids)
- VTE prophylaxis
- Carbohydrate load (2hr preop)
- Confirm stoma marking
- Confirm CCM aware of ostomy

## OR

- Goal-directed fluid administration
  - 2L total in OR
  - 3L total/first 24hrs
- Open procedures: No epidural

## Postop Day 0

- No sitting
  - Order Sign over bed “No sitting”
- Goal-directed fluid administration
- OOB (Dangling not compliant)
- Diet
  - Low Residue diet
  - Ensure Supplements
- Teaching
- Gum, Mag & Entereg
- Pain Management
  - PCA
  - Tylenol 1gm q6 ATC
  - Gabapentin 300mg TID
  - Oxycodone once at night
  - Tramadol PRN
  - Resume all baseline pain meds
- Home Medications
  - Resume all home medications
  - No therapeutic anti-coagiation
  - Diabetes medicines
    - \* Prefer to resume all
    - \* Sliding-scale insulin if needed
- *Wound Ostomy teaching*
- *CCM for Home Health for stoma care*

### Postop Day 1

- Labs: K+ and CBC only
- d/c “pre” plan & colon visit
- Heparin lock IV
- Out of bed > 6 hrs
- Diet: low residue
- Pain management
  - 1) d/c PCA
  - 2) Tylenol
  - 3) Gabapentin 300 tid
  - 4) Tramadol
  - 5) Home meds
  - 6) Oxy if lots of pain

### Postop Day 2

- No IV fluids unless indicated
- No labs unless indicated
- OOB >6 hrs
- No PT unless going to rehab or SNF (ask Hill first)
- Pain management
  - d/c PCA (unless laparotomy)
- *Wound Ostomy Teaching*
- Foley voiding challenge
  - 250mL saline into foley -> pull
  - Must void within 2 hours
- Discharge planning
  - Consider early D/C
- Otherwise plan
  - Patient
  - Nursing
  - *CCM for home health for stoma care*
- Afternoon rounds
  - 1) Possible home today!
  - 2) Check patient 2-3PM
  - 3) Ambulation 2x's by PM
  - 4) Check for dehydration
  - 5) Patient education

- 6) Give estimated date of discharge

**Postop Day 3-5** - IVF - Consider bolus of IV fluids - Consider maintenance IV if we feel won't resolve soon - Labs - No labs unless indicated - Consider ordering QOD Chem7 if prolonged ileus - OOB >6 hrs - No PT unless likely to go to rehab or SNIF (ask Hill first) - Pain management - Same as POD#1 - Afternoon rounds 1) Possible home today! 2) Check patient 2-3PM 3) Ambulation 2x's by PM 4) Check for dehydration 5) Patient education 6) Give estimated date of discharge

## Chapter 13

# Esophagectomy Postop

### ICU Care

All patients admitted to STICU with Surgical Critical Care Consultation. Surgical critical care will need a phone call immediately after surgery at 6-0366. Listed: Dr Salo, CMC-GI Surgical Oncology, CMC-Surgical Critical Care. Diabetic patients (on insulin preop): Consult Kelli Dunn.

### Ward Care

Once stable, patients are transferred to 6T. If a 6T bed is not available, please notify Dr Salo. Historically, over 95% of patients are transferred to 6T after leaving the ICU.

### Neuro

Multimodal pain control:

- gabapentin (300mg tid liquid via Jejunostomy)
- Tylenol (1000mg q6hrs as pediatric liquid via Jejunostomy).
- PCA with subsequent conversion to oxycodone elixir via Jejunostomy.
- No ketorolac (Toradol) given risk of anastomotic failure<sup>1</sup>
- Home anxiolytics are generally administered at half the home dose

### Cardiovascular

Postoperative atrial fibrillation is a common occurrence (20%) after esophagectomy, with the risk increasing in older patients. For patients over age 70, half of patients will develop atrial fibrillation in the postoperative period.

For prevention of atrial fibrillation, beta blockade is used. For patients receiving beta blockers prior to surgery, continuation of beta blockade is recommended. For others, patients are given metoprolol 2.5 mg IV q6hrs which can be titrated up to 10mg IV q6hrs as needed. See STS Guidelines.<sup>2</sup>

Home anti-hypertensives are usually held in order to allow beta-blockade. Patients who have elevated blood pressures once they are adequately beta-blocked (eg HR 60-70) are usually restarted on their home anti-hypertensives. *Home anti-hypertensives are not routinely restarted postoperatively*

### Respiratory

Chest tubes generally consist of a 28Fr Blake drain placed into the right chest. This is placed to water seal when output is less than 200mL/day and there is no leak visible in the Pleurevac container. Chest tubes are usually removed once output is less than 150mL/day and drainage is clear without evidence of chyle (milky appearance). A chest x-ray is obtained after removing a chest tube to look for a pneumothorax.

**Gastrointestinal** All patients receive pantoprazole 40mg IV daily and metoclopramide 5-10mg IV q 6hrs

### Nutrition

All esophagectomy patients receive a feeding jejunostomy at the time of operation. In the immediate post-operative period, patients receive Osmolite 1.5 starting the day of surgery once they are off pressors. Tube feeds are started at 20mL/hour until flatus and then advanced at 10mL per hour every 8 hours, up to a goal of 60mL per hour (x24 hours). Patients who are on tube feeds prior to surgery are generally restarted on their home tube feed formula.

Patients who do not tolerate Osmolite are switched to Vital 1.5, which is pre-hydrolyzed. In order to allow enough time for switching of tube feedings, patients are generally switched from Vital to their home tube feeding formula 3-4 days prior to discharge. This is typically done when they are transferred out of the ICU.

Obese patients (BMI>30) are started on Promote at 20mL/hour and increased to a goal of 60mL/hour. Promote contains a more protein relative to carbohydrates. An alternative is Vital High Protein, which is similar to Promote but using hydrolyzed proteins.

Diarrhea in patients on tube feeds (especially nocturnal diarrhea) needs to be addressed. Despite STICU guidelines for nutrition in trauma patients, diarrhea (especially night-time diarrhea) is justification for alteration in tube feeds. See Diarrhea and Jejunostomy Feeding

### Diabetic Patients

Patients who require EndoTool in the ICU will need an endocrinology consult. See also Jejunostomy Feedings with Diabetes

### 'Free Water'

In addition to tube feeds, most patients will receive 'free' water flushes through the jejunostomy. This is typically done as 240mL four times per day (for a total of 32oz)



### *Nasogastric Tubes*

A silicone nasogastric tube (Covidien Salem Sump) is placed in all patients during surgery and the position confirmed by ultrasound intraoperatively. Tubes are positioned so that all 4 dots are outside. Gastric emptying is evaluated with upper GI prior to NG tube removal. Once extubated, upper GI is typically performed on the 2nd through 4th postoperative day. Conversely, patients who are intubated will keep their nasogastric tube until extubated. Radiology ordered as “upper GI Series”. In the comments section please add “IsoVue through NG tube. Halo Dr Salo for study”. See Evaluation of GI Function

### *Drains*

- JP1: 19Fr Blake drain in left pleura. The exit site the most lateral drain and is secured with a blue suture
- JP2: 19Fr Blake drain in right pleura. The exit site is medial to JP1 and is secured with a black suture
- JP3: 19 Fr Blake drain in abdomen. If used, the exit site is most medial and is secured with a blue suture
- JP4: 15Fr Blake drain in neck (for cervical incision)
- JP5: 15Fr Blake drain in subcutaneous tissue of incision

### *Evaluation for Anastomotic leak*

Drain amylase is an inexpensive, specific, and relatively sensitive test for anastomotic leak. Fluid from JP2 (right chest) is sent for “Body Fluid Amylase” starting on postoperative day #4 and continued until postoperative day 9. Drain amylase over 400IU/ml is considered positive and prompts a CT esophagram for confirmation. See also Evaluation for Anastomotic Leak

**Renal** Total fluids (IV + tube feeds) are generally run at 75mL/hour. Foley catheter is removed on the first or second day after surgery. Some patients will need diuresis on the 3rd or 4th postoperative day

**Heme** All patients require VTE prophylaxis with Lovenox or heparin SQ.

Preoperative anti-platelet agents are started on the first (aspirin) or second (Plavix) postoperative day if there is no excessive bleeding from the chest tube or JP drains. Patients on preoperative anticoagulation are transitioned to therapeutic Lovenox on the second postoperative day if no signs of bleeding.

**ID** Prophylactic Cefazolin and Flagyl are administered for 24 hours and stopped



## Chapter 14

# Esophagectomy Ward

### Ward Care

Once stable, patients are transferred to 6T. If a 6T bed is not available, please notify Dr Salo. Historically, over 95% of patients are transferred to 6T after leaving the ICU.

### 14.1 Anti-Hypertensives

Patients are transitioned to enteral metoprolol once they are transferred to the ward. Patients on IV metoprolol at 2.5mg IV q6 hours are started on 25mg enteral bid, while patients receiving 5mg IV q6 hours are started on 50mg bid. For patients who are not taking medicines by mouth, liquid metoprolol can be ordered while an inpatient. (Liquid metoprolol is not available as a home medicine.)

Home anti-hypertensives are usually held in order to allow beta-blockade. Patients who have elevated blood pressures once they are adequately beta-blocked (eg HR 60-70) are usually restarted on their home anti-hypertensives. *Home anti-hypertensives are not routinely restarted postoperatively.* Home antihypertensives are restarted on a selective basis

### 14.2 Chest tubes

Chest tubes generally consist of a 28Fr Blake drain placed into the right chest. This is placed to water seal when output is less than 200mL/day and there is no leak visible in the Pleurevac container. Chest tubes are usually removed once output is less than 150mL/day and drainage is clear without evidence of chyle

(milky appearance). A chest x-ray is obtained after removing a chest tube to look for a pneumothorax.

### 14.3 GI Medicines

All patients receive pantoprazole 40mg IV daily and metoclopramide 5-10mg IV q 6hrs. This is later switched to enteric PPI and reglan. For patients <age 75, remeron is added as 15mg enteral qhs.

### 14.4 Evaluation for leak

Experience suggests that the median time to the diagnosis of leak is 7 day after surgery. Currently the risk of anastomotic leak at CMC after transthoracic esophagectomy is 2%. The most sensitive test for the diagnosis of leak is CT esophagram (see below). Based upon institutional experience, patients are divided into low risk of leak vs high risk for leak based upon drain amylase level and white blood cell count. Patients with a normal drain amylase (400IU/ml) between postoperative day 4 and day 7 AND white blood cell count less than 12 are considered low risk. Patients with either an elevated drain amylase or WBC greater than 12 are evaluated with CT esophagram. In a group of 100 patients, several were found to have elevated drain amylase in the first four days (up to 2000Iu/ml) which subsequently declined, and no evidence of leak was found

#### 14.4.1 Drain Amylase

JP2 is placed into the right pleura, passes through the hiatus, and is brought out through a trocar site in the medial left upper quadrant. Drain amylase from JP2 is tested beginning on postoperative day #4 until discharge or postoperative day #9. JP2 is generally removed prior to patient discharge. JP1 is placed in the left pleura and is generally not tested for amylase.

#### 14.4.2 CT esophagram

This is the most sensitive study for the detection of anastomotic leak. In order to obtain sufficient sensitivity, it requires a pre-contrast scan and the administration of contrast into the esophagus. The need for a pre-contrast scan means that the presence of remnant barium in the esophagus (from a Modified Barium Swallow) makes it more difficult to interpret the scan and should be avoided. Awake patients can drink the contrast. Patients with an NG tube present at the time of the study generally will have contrast administered through their NG tube. Almost all esophagectomy patients will have a Covidien silicone Salem

Sump 18Fr tube in place. This has four marks on the tube at 45, 55, 65, and 75cm. The tube is typically positioned with the 4th mark at the nares, which means that the tip is 45cm from the nares and is usually within the gastric conduit AND below the anastomosis. The NG tube also contains side holes which extend 8.5cm above the tip of the tube. The optimal study is done with the NG tube withdrawn so that there are two side-holes above the anastomosis, which means that the tip is approximately 6cm below the anastomosis. A scout CT will be performed, and the radiologist will determine how far back the NG tube needs to be withdrawn. This is communicated to the CT technician, who asks the nurse (or physician) to withdraw the tube the calculated amount (making a note of the starting position of the tube relative to the four marks). This should keep the tip below the level of the anastomosis, so that after the CT scan, the NG tube can be (blindly) advanced back to its original position (at approximately 45cm from the nares).

Patients who clinically deteriorate prior to post-operative day 7 in whom there is high suspicion for a leak (fevers, pleural effusion on chest X-ray, elevated JP amylase, respiratory failure) will undergo a CT esophagram earlier than post-operative day 7.

## 14.5 Anastomotic Leak Treatment

If a patient is demonstrated to have a leak, they are made NPO AND will have any pleural effusion treated with a pigtail catheter. Conservative management will generally be successful for most patients with leaks provided 1) there is no evidence of conduit necrosis 2) nutrition is optimized and 3) empyema is treated. Patients with leaks may even need a decortication, which emphasizes the importance of CT scan in the sick post-operative esophagectomy patient, so that an empyema can be diagnosed and treated. Patients with leaks who show signs of systemic illness may need to be considered for an intraluminal stent. Patients who are profoundly ill need to be evaluated for gastric necrosis with upper endoscopy.

### **Nutrition**

All esophagectomy patients receive a feeding jejunostomy at the time of operation. In the immediate post-operative period, patients receive Osmolite 1.5 starting the day of surgery once they are off pressors. Tube feeds are started at 20mL/hour until flatus and then advanced at 10mL per hour every 8 hours, up to a goal of 60mL per hour (x24 hours). Patients who are on tube feeds prior to surgery are generally restarted on their home tube feed formula.

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Diarrhea in patients on tube feeds (especially nocturnal diarrhea) needs to be addressed. Despite STICU guidelines for nutrition in trauma patients, diarrhea (especially night-time diarrhea) is justification for alteration in tube feeds. See Diarrhea and Jejunostomy Feeding

#### *Diabetic Patients*

Patients on tube feeds are typically started on continuous (around-the-clock) tube feedings, and subsequently changed to a nocturnal regimen (typically 6pm to 10am). See Jejunostomy Feeds in Diabetic Patients

#### *‘Free Water’*

In addition to tube feeds, most patients will receive ‘free’ water flushes through the jejunostomy. This is typically done as 240mL four times per day (for a total of 32oz)

#### *Nasogastric Tubes*

A silicone nasogastric tube (Covidien Salem Sump) is placed in all patients during surgery and the position confirmed by ultrasound intraoperatively. Tubes are positioned so that all 4 dots are outside. Gastric emptying is evaluated with upper GI prior to NG tube removal. Once extubated, upper GI is typically performed on the 2nd through 4th postoperative day. Conversely, patients who are intubated will keep their nasogastric tube until extubated.

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**Heme** All patients require VTE prophylaxis with Lovenox or heparin SQ.

Preoperative anti-platelet agents are started on the first (aspirin) or second (Plavix) postoperative day if there is no excessive bleeding from the chest tube or JP drains. Patients on preoperative anticoagulation are transitioned to therapeutic Lovenox on the second postoperative day if no signs of bleeding.

**ID** Prophylactic Cefazolin and Flagyl are administered for 24 hours and stopped

Discharge Medicines Patients after esophagectomy typically go home with the following medicines:

Proton pump inhibitors (will continue for 2 years) Oxycodone elixir via feeding tube. Current STOP guidelines dictate that patients receive no more than a 7 day supply of opioids at discharge Reglan 10mg po qid (will stop at 6 weeks post-op) Remeron 15mg qhs (will continue for 3 months post-op) Tylenol 1000mg q6 hours as elixir (pediatric form) Gabapentin 300mg as liquid tid x 14 days Metoprolol if started postoperatively (most patients). If not on beta blockers preoperatively, this will be cut in half at the first visit and stopped at the second postoperative visit. It is important to limit the use of medicines via jejunostomy in order to lower the risk of clogging the feeding tube. While liquid metoprolol is available for inpatients, it is not available from outpatient pharmacies.





## Chapter 15

# Jejunostomy Feedings

Due to the osmotic load, jejunostomy feedings are given via enteral (Kangaroo) pump rather than bolus feeding. Feedings are generally begun as continuous (around-the-clock) and are then transitioned to nocturnal (generally 6pm to 10am) prior to discharge.

Jejunostomy feedings carry a small but significant risk (~1%) of small bowel necrosis, as evidenced by the findings of pneumatosis on CT scan and in some cases small bowel necrosis and perforation. The existing literature would suggest that the early symptoms associated with small bowel necrosis are abdominal distension. As a result, patients on jejunostomy tube feeding need to be carefully monitored for distension, and tube feeds held if distension develops. (Taylor et al., 2014) Tube feeds are generally started at 30mL/hour in the immediate postoperative patients. In patients who are awake and in whom it is possible to determine whether or not there are issues of tube feed intolerance, the rate of tube feedings is increased 10mL/hour every 12 hours to a goal of 60mL/hour for women and 75mL/hour for men. In patients who are intubated/sedated, the advancement of tube feeds is individualized, and decisions are made on a daily basis on rounds.

For patients who receive tube feedings preoperatively, the same formula is generally used after surgery.

For patients with BMI less than 35, Osmolite 1.5 is used as a starting formula. If Osmolite is not tolerated due to diarrhea, Vital 1.5 can be used. For patients with BMI greater than 35, Promote is used as a starting formula, as it contains a lower amount of carbohydrates than Osmolite 1.5. Patients who are intolerant to Promote can be switched to Vital High Protein.

Patients less than 150 pounds receive 4 cans of tube feedings, administered at 60mL/hour x 16 hours (6pm to 10am). Patients greater than 150 pounds receive 5 cans of tube feedings, administered at 75mL/hour x 16 hours (6pm to 10am). Patients with BMI >35 receive 5 cans of Promote (or Vital High Protein).

Glucerna is formulated with carbohydrates with low glycemic index, which is particularly helpful for bolus feeding (via gastrostomy) or patients who are eating. Glycemic index is less important in patients on continuous tube feeds (eg via jejunostomy). In addition, the high fiber context of Glucerna may make this formula more prone to causing clogging of jejunostomy tubes. Free Water Flushes

Once patients are transferred to the ward, free water via jejunostomy should be ordered as 240mL via jejunostomy qid. This is entered under Flush Enteral Tube for Free Water Requirements. Diarrhea with Jejunostomy Feedings

Patients who experience diarrhea during jejunostomy administration (especially diarrhea at night) will need this addressed. Several steps

- Send stool for C Diff
- Consider changing tube feedings to a more easily digestible formula. Patients on Osmolite 1.5 can be changed to Vital 1.5. Patients on Vital 1.5 can be changed to Vivonex. Patients on Promote can be changed to Vital High Protein
- Stop tube feedings for 2-4 hours to allow diarrhea to resolve
- restart tube feeds at a lower rate (eg 20mL/hour lower than the prior rate).
- Bannatrol can be given to patients taking an oral diet. Bannatrol is NOT given via jejunostomy tube to avoid clogging.
- Lomotil is generally used as a last resort.

## 15.1 Med Administration via J-Tubes

Patients who have difficulty with dysphagia or complete esophageal obstruction will need to have their medicines administered via jejunostomy tube. The process of designing a medicine regimen which can safely be administered via enteral tube can be a challenge and may require a consultation with the hospital pharmacist. One the other hand, the financial consequences of a clogged feeding tube are substantial. **Flomax is never given via jejunostomy tubes due to risk of clogging.**

Several common medicines are available in liquid form:

- Acetaminophen (pediatric formulation)
- Gabapentin
- Oxycodone
- Hydrocodone + acetaminophen
- Reglan

## 15.2 Occluded Jejunostomy Tubes

Jejunostomy tubes which are refractory to the usual non-invasive means (warm water, Coca-Cola) will need to be changed over a wire in Interventional Radiology.

## 15.3 Jejunostomy + Diabetes

### Inpt – Jejunostomy ~ Diabetes

Patients on tube feeds are typically started on continuous (around-the-clock) tube feedings, and subsequently changed to a nocturnal regimen (typically 6pm to 10am). The diabetic management for these patients differs depending upon their tube feeding regimen:

#### **Diabetics on Continuous Tube Feeds.**

Non-insulin diabetic patients are generally initially given Osmolite 1.5 as it provides higher caloric density and does not contain fiber, which tends to clog the feeding tubes. Diabetic patients requiring insulin can be initially trialed on Osmolite 1.5 but are changed to Promote or Glucerna 1.5 if their blood sugars prove difficult to control (as evidence by either the need for EndoTool or requiring a q6 hour regimen of insulin N + insulin R).

Diabetic patients who need insulin while receiving tube feedings are typically treated initially with continuous tube feedings and around-the-clock insulin. Patients with large insulin requirements may need hourly intravenous insulin (with dosages calculated via EndoTool). Once their insulin requirements are stabilized, they are transitioned to a q6 hour regimen consisting of N and R insulin, typically twice as many units of N insulin as R (for instance, 6Units of N + 3 Units of R insulin every six hours). Patients who require EndoTool or a q6hr regimen need an endocrinology consultation with Dr Kelli Dunn to assist in diabetic management.

#### **Diabetics on Nocturnal Tube Feeds**

Most patients are transitioned from continuous tube feeds to nocturnal prior to discharge. Diabetic patients on nocturnal tube feedings typically receive tube feeding from 6pm to 10am and receive insulin at initiation of tube feeds (6pm) and again at 6 hours later (midnight). A typical regimen might be 18U of 70/30 at 1800 and 18U of 70/30 at MN. An alternative might be 12U NPH + 6U Regular at 1800 and 16U NPH and 8U Regular insulin at MN. Because it can take several days to determine the correct insulin regimen, diabetic patients receiving jejunostomy feedings are cycled as early in their hospital course as possible to avoid delaying discharge for blood sugar management.

Diabetic patients receiving insulin will need careful coordination of tube feeding and insulin administration when they are being transitioned from continuous to

nocturnal tube feeds. Patients on continuous tube feeds may receive insulin on a q6 hour schedule, while those receiving nocturnal tube feeds receive insulin at 1800 and MN. When patients on continuous tube feeds are transitioned, the tube feeds are stopped at 10am, to be restarted at 6pm that evening. It is critical that as soon as the tube feeds are stopped at 10am, that the q6 hour insulin as stopped as well.

In either case it is critical that if tube feedings are stopped, standing insulin administration (either q6 hour OR 1800 and MN) be stopped as well. In these cases, sliding scale insulin is generally continued.

Day	Time	Tube Feeds	Insulin
SUN	MN	60mL/hr	8N+4R
Mon	6am	60mL/hr	8N+R
Mon	Noon	Stop	None
Mon	6pm	75mL/hr	16N + 8R
Mon	MN	75mL/hr	16N+ 8R
Tues	10am	Stop	None
Tues	6pm	75mL/hr	16N+ 8R
Tues	MN	75mL/hr	16N+ 8R
Weds	10am	Stop	None

In this example, the patient was receiving  $8N + 4R$  every 6 hours, so total insulin units per day is  $(8+4) \times 4 = 48$  units. Because the carbohydrate load of 60mL/hour  $\times$  24 is roughly equivalent to 75mL/hour  $\times$  16 hours, the total insulin administered is roughly the same. When converted to nocturnal dosing, the patient now received  $16+8 = 24$  units twice (6pm and MN) = 48U. In practice, it may be wiser to begin by adjusting the dose down a little, to perhaps 14U N and 7U R for the first night.

**Hill OR**



## Chapter 16

# Colorectal Cases - Hill

### **Pre-op holding**

Please make sure that these ordersets have been placed prior to day of surgery. If inpatient consult, place these orders the morning of surgery in order to minimize confusion.

ADULT SURG Colorectal ERAS MPP Hill This is what has all of the main ERAS components. Tylenol, gabapentin, Decadron, Alvimopan and heparin are all given in pre-op holding

ADULT STANDING Antimicrobial colorectal In general, I will give Ancef to patients with almost all patients with allergies to PCN. They have to remember the “severe” reaction. If it is a severe reaction, please use the second line ABX listed in the power plan.

Type and Screen are not typically needed for colectomy. They will have an antibody screen from office. If antibodies present, then d/w attending

### **Intra-op**

Positioning: I typically like to position myself.

Right sided=supine; Left sided=lithotomy; All laparoscopic colon cases will have their arms tucked with a chest tape strap.

NG/OG tubes Not needed. I will have anesthesia place if we have gastric distension. We give a multiple PO meds prior.

Review anesthesia fluid management during time out. 1.6L max volume, urine output not an accurate indicator.





**Salo OR**



## Chapter 17

# Central Venous Port (IJ)

### Room Prep

- Slider bed (Skytron 3600B) with head section
- C-Arm
  - Radiology technician alerted to need for C-Arm
  - Will need lead and thyroid shields for everyone in room
- BK Ultrasound with hockey-stick probe near patient's RIGHT SHOULDER

### Instruments - Minor instrument pan

### Disposables/Meds

- Confirm choice of port with surgeon. Usual options
  - Bard PowerPort VUE with 8Fr attachable catheter (1708062)
  - Bard PowerPort slim Implanted port (for patients with low BMI)
    - Heparin 5mL of 1000U/ml labeled as “1000 U/ml” -Heparin 5mL of 1000U/ml + 45mL saline labeled as “100 U/ml”
  - Local
    - \* If general anesthesia: Marcaine 0.5% with epinephrine
    - \* If MAC: Xylocains 1% with epinephrine
- 1000 drape x3 AND blue paper drapes 4 packs of 2 each = 8 total
- Suture
  - 3-0 Prolene RB-1 double-arm
  - 3-0 Vicryl SH
  - 4-0 Monocryl PS2

**Position**

- Supine with left arm tucked, right arm on armboard at side.
  - Right arm on armboard in case needed by anesthetist
  - NO shoulder roll
- Foley catheter: usually NOT required – *check with surgeon*
- Lower body Bair Hugger from abdomen to feet with ONE layer of blankets on top of Bair Hugger. Velcro strap on thighs.

**Prep**

Chloroprep: RIGHT chest, neck to chin and earlobe, shoulder to include deltopectoral groove.

**Drape**

1000 plastic drapes outline the sterile field for the port. The skin is stretched to avoid a gap between drape and skin. Allow access to the right sternocleidomastoid, right deltopectoral groove, and sternal notch. Blue paper drapes on top of 1000 drapes Transverse drape reversed head-to-foot. Ioban around edges of port field. Skin over SCM is left without Ioban to facilitate ultrasound

**Preop evaluation** Allergies Blood thinners or anti-platelet agents History of prior central venous lines or ports History of neck surgery

**Operation***Reverse Trendelenburg*

Port pocket is constructed 1cm below and parallel to clavicle 3cm long. It is essential that there is no bleeding in the pocket (to avoid a port pocket hematoma).

*Trendelenburg*

Right internal jugular vein is identified and its course cephalad-caudad marked on the skin.

Finder needle (22Ga) *OR* micropuncture kit passed into IJ. The needle should enter the vein directly beneath the ultrasound probe.

Skin anesthetized and transverse 8mm counter-incision made at needle entry site

*Respiration held by anesthesia*

16Ga needle passed into IJ under sono. It is essential that the vein is scanned up and down by ‘rocking’ the probe to visualize the tip of the needle as it passes inferior.

J Wire passed through 16Ga needle and needle withdrawn

*Anesthesia resumes respirations*

Ultrasound used to confirm presence of wire within the vein by scanning up and down.

*Level bed*

Fluoroscopy used to confirm position of wire. Dilator and sheath inserted under fluoroscopic visualization ('live'). C-arm backed away off field.

Tunneler connected to tubing on the end with small numbers. Confirm that the collar is in place and place hemostat on the end of the tubing with large numbers (to avoid allowing the collar to fall off). Tunneler bent into a curve to avoid injury to the carotid artery.

Catheter tunneled from port pocket to counter-incision over SCM. The tunneler path describes a gentle arc to avoid kinking the catheter. 1 cm of catheter near tunneler trimmed.

Catheter placed through dilator into central circulation. Most of the catheter is inserted. The catheter will generally not cause arrhythmias.

C-arm brought back onto field.

Peel-away sheath split and removed.

Traction on catheter from the port pocket is used to position catheter approximately 3cm below carina. It may be necessary to 'orbit' the C-arm if the catheter overlies the spine.

Port pocket is measured and catheter trimmed (after sliding collar superior) and attached to port. It is essential that the catheter come to rest within 1mm of the port before locking the collar in place. In order to avoid pulling on the catheter (and changing the position of the catheter tip) the port is rotated (not the catheter). Collar is locked in place.

Port accessed with straight Huber needle with 100U/ml heparinized saline. Blood is withdrawn into port. Needle is left in the port and the syringe detached. Syringe with concentrated flush (1000U/ml) is attached to the needle and the port flushed (without aspiration of blood). Syringe and needle are removed.

Port sutured to the underlying pectoralis fascia with 2 sutures of 3-0 Prolene, one forehand and one backhand. Sutures are tied and cut.

The port pocket irrigated and the incision closed with subcutaneous 3-0 Vicryl followed by subcuticular 4-0 Monocryl. The skin is dressed with Dermabond.

**Postop Orders**

CXR in recovery to confirm central line placement



# Chapter 18

## Lap Jejunostomy

### Room Prep

- EGD cart near patient's LEFT SHOULDER (with ADULT EGD scope)
- If central venous port is placed at the same time:
- Slider bed (Skytron 3600B) with head section
- Radiology technician alerted to need for C-Arm
- BK Ultrasound with hockey-stick probe near patient's RIGHT SHOULDER

### Instruments

- 5mm 30 degree scope AND 5mm 0 degree scope
- SRI laparoscopic Pan
- Salo laparoscopic instruments

### Disposables/Meds

- Veress needle (with 10mL syringe and saline)
- 5mm Z-thread optical port (3 on table, 2 more in room)
- Transverse drape AND laparoscopy drape
- Confirm choice of port with surgeon. Usual options
  - Bard PowerPort 8Fr xx8062
  - Bard PowerPort 8Fr xx8000 (low profile)
- Heparinized saline: 100U/ml (dilute) and 1000U/ml (concentrated)
- 1000 drape x3 AND blue paper drapes 4 packs of 2 each = 8 total
- Micropuncture kit available/not open (from Anesthesia)
- Jejunostomy tube: MIC 0301-14

- Silk 2-0 on RB-1 needle (on Surgical Oncology suture cart)

### Position

- Supine with left arm tucked, right arm on armboard at side.
- Foley catheter: Usually required – *check with surgeon*
- Lower body Bair Hugger on thighs. ONE layer of blankets on top of Bair Hugger. Velcro strap on thighs. NO PILLOW UNDERNEATH LEGS.

### Prep

Chloroprep (two sticks) of abdomen (need to keep pubis in field, as well as right anterior superior iliac spine), both costal margins.

If port: RIGHT chest, neck to chin and earlobe, shoulder to include deltopectoral groove

### Drape

If central venous port: Perimeter of field draped with 1000 (clear adhesive) drapes. Four 1000 drapes around port site:

- Medial border: From Angle of Louis superiorly along midline to chin.
- Superior border: Inferior to jaw (to allow access to right internal jugular vein and SCM)
- Laterally: From inferior to ear down to right shoulder
- Inferior: From lateral shoulder medially to Angle of Louis

Abdomen: Two 1000 drapes used inferiorly keeping pubis and right anterior inferior iliac spine in field. This is critical as the far inferior/lateral RLQ needs to be in the field for optimal port placement.

Six Blue Paper Drapes around perimeter of field (on top of 1000 drapes)

If central venous port: Transverse sheet TURNED HEAD-TO-FOOT turned at an angle to keep deltopectoral groove and SCM within the field

Laparoscopy drape skewed to inferior and right to keep pubis and right ASIS in the field.

Turn on Bair Hugger only AFTER drapes in place

### Indications

Laparoscopic jejunostomy is used for enteral nutrition in patients prior to planned (or possible) esophagectomy or gastrectomy or those for whom the stomach is otherwise not available (ie after esophagectomy or gastrectomy). Patients with metastatic esophageal cancer who need enteral access are generally treated with a gastrostomy, which does not require feeding with a pump



Preop (Resident) Preop orderset: search for “Jejunostomy”

Review Clinical Information (Resident) Review staging scans (especially PET scan) to identify suspicious areas on imaging which need to be investigated at the time of laparoscopy Outpatient anticoagulation use (warfarin, Xaralto, aspirin, Plavix) Review dietitian’s recommendations (how many cans of feeding per day?) If patient is scheduled for central venous port Confirm that a port has not already been placed Prior history of central venous lines? Confirm location of port placement with surgeon (left vs right)

### **Operation**

If a central venous port is placed, the port is performed first. See IJ Port

Abdominal access is obtained in one of two ways:

Infraumbilical approach using modified Hasson technique. If the peritoneum is not easily entered, a Veress needle is used to insufflate, followed by incision of the fascia with a 15 blade, and a 5mm optical port (Applied Medical Kii Fios First Entry Z-Thread Trocar) Veress needle inserted in LEFT upper quadrant just inferior to costal margin. Abdominal entry with 5mm optical Z-Thread port. 5mm port in right upper quadrant, 5mm port in RLQ just lateral to rectus, 5mm camera port between RUQ and RLQ ports

The transverse colon is now elevated (using the umbilical port, if used) and the ligament of Treitz is identified. The proximal bowel is arranged in a “C” configuration to confirm the proximal and distal ends of bowel.

A site for placement of the jejunostomy is selected on the skin, left lateral and just superior to the umbilicus. A site is selected on the bowel in the most proximal site on the jejunostomy selected which would allow for placement of the jejunostomy without tension, but at least 20cm from ligament of Treitz.

The proximal jejunum is sutured to the anterior abdominal wall with 2-0 silk. This is usually done with a 9 suture which is introduced into the abdomen with a needle driver “Korean Style” or “Paraguayan Style.” Two cm distal to this suture, a diamond of sutures is placed around the proposed tube site, and one suture placed distal to avoid torsion. The final arrangement of sutures is one proximal and one distal and 4-6 sutures around the tube. All sutures were marked with hemostatic clips to facilitate replacement of the tube via fluoroscopy should the tube become dislodged.

Using Seldinger technique, a 16Fr Cook catheter introducer kit is placed within the jejunum.

A MIC 14Fr jejunostomy tube (0301-14) with the tabs trimmed with a scalpel, is inserted through the sheath and positioned in the jejunum. The tube was secured with a suture of 0 silk.

If a balloon tube is used, an 18Fr Cook dilater and sheath is used, followed by a MIC 14Fr jejunostomy tube (0200-14) and the balloon inflated with 7mL of sterile WATER.

The tube is secured with 0 silk and dressed with a BioPatch and a Tagederm dressing.

The abdomen is desufflated and the port sites closed with 4-0 Monocryl, followed by dermabond.

After dressings are applied, a Lopez valve is attached with the long Christmas-tree end placed into the jejunostomy tube.

Endoscopy The scope is set up:

Suction and aspiration valves inserted and working Suction tubing attached Biopsy valve attached and not leaking Cart set up for recording by powering on the Stryker SDC digital capture box A bite block is used and the scope lubricated. A neonatal scope may be necessary in patients with a tight stricture. Important findings to record:

Level in cm from the incisors, of the most proximal area of Barrett's esophagus.  
Level in cm from the incisors of the GE junction Appearance of the GE junction on retroflexed view. Extent of invasion of the tumor into the cardia or fundus.  
The scope is withdrawn and the hypopharynx suctioned. The liquid from the 'First Step' disinfectant is suctioned through the scope, followed by water.

# Chapter 19

## Lap Gastrostomy

### Room Prep

- EGD cart near patient's LEFT SHOULDER (with NEONATAL EGD scope)
- If central venous port is placed at the same time:
  - Slider bed (Skytron 3600B) with head section
  - Radiology technician alerted to need for C-Arm
  - BK Ultrasound with hockey-stick probe near patient's RIGHT SHOULDER

### Instruments

- 5mm 30 degree scope AND 5mm 0 degree scope
- SRI laparoscopic Pan (available)

### Disposables/Meds

- Veress needle (with 10mL syringe and saline)
- 5mm Z-thread optical port (3 more in room)
- Transverse drape AND laparoscopy drape
- Confirm choice of port with surgeon. Usual options
  - Bard PowerPort 8Fr xx8062
  - Bard PowerPort 8Fr xx8000 (low profile)
- Heparinized saline: 100U/ml (dilute) and 1000U/ml (concentrated)
- 1000 drape x4 AND blue plastic adhesive drapes 4 packs of 2 each = 8 total
- Micropuncture kit available/not open (from Anesthesia)

- 20Fr Laparoscopic gastrostomy kit in room/not open
- 16Fr MIC gastrostomy tube in room/not open
- Gastrostomy 20Fr Pull PEG (in vending machine)
- GI Anchors (“T-fasteners”) in room/not open

### Endoscope Setup – Neonatal EGD scope

- Valves attached and working (suction/aspiration/biopsy cap)
- Suction tubing attached
- Connect water bottle to left-hand port
- Gauze sponges, lubricant, plastic “tray” from gauze filled with water
- **Yellow** (small) bite block
- “First step” sanitizer

### Anesthesia

- ET Tube taped to left. Head turned to the left on donut
- No EKG electrodes on anterior right chest

### Position

- Supine with left arm tucked, right arm on armboard at side.
- Foley catheter: usually NOT required – *check with surgeon*
- Lower body Bair Hugger on thighs. ONE layer of blankets on top of Bair Hugger. Velcro strap on thighs. NO PILLOW UNDERNEATH LEGS.

### Prep

Chloroprep (two sticks) of abdomen (need to keep pubis in field, as well as right anterior superior iliac spine), both costal margins.

If port: RIGHT chest, neck to chin and earlobe, shoulder to include deltopectoral groove

### Indications

Laparoscopic gastrostomy is used in patients with esophageal obstruction . Gastrostomy feedings are much easier than jejunostomy, as they can be administered via syringe or gravity bag. By contrast, jejunostomy feedings require administration via pump. Gastrostomy is usually done as an outpatient unless there are concerns for refeeding.

### Review Clinical Information (Resident)

Review staging scans (especially PET scan) to identify suspicious areas on imaging which need to be investigated at the time of laparoscopy Outpatient anticoagulation use (warfarin, Xaralto, aspirin, Plavix) Review dietitian’s recommendations (how many cartons per day?)

## Drape

If central venous port: Perimeter of field draped with 1000 (clear adhesive) drapes. Four 1000 drapes around port site:

- Medial border: From Angle of Louis superiorly along midline to chin.
- Superior border: Inferior to jaw (to allow access to right internal jugular vein and SCM)
- Laterally: From inferior to ear down to right shoulder
- Inferior: From lateral shoulder medially to Angle of Louis

Abdomen: Blue adhesive drapes used inferiorly keeping pubis and right anterior inferior iliac spine in field. This is critical as the far inferior/lateral RLQ needs to be in the field for optimal port placement if the patient needs a jejunostomy.

Six Blue Adhesive Drapes around perimeter of field (on top of 1000 drapes)

If central venous port: Transverse sheet TURNED HEAD-TO-FOOT turned at an angle to keep deltopectoral groove and SCM within the field

Laparoscopy drape skewed to inferior and right to keep pubis and right ASIS in the field.

Turn on Bair Hugger only AFTER drapes in place

## Operation

If a central venous port is placed, the port is performed first. See IJ Port

**Abdominal access** is obtained in one of two ways:

- Veress needle inserted in left upper quadrant just inferior to costal margin. Abdominal entry with 5mm optical port at lateral border of rectus just superior to umbilicus
- Infraumbilical approach using modified Hasson technique. If the peritoneum is not easily entered, a Veress needle is used to insufflate, followed by incision of the fascia with a 15 blade, and a 5mm optical port

The insufflation pressure is decreased to 4mmHg and the abdomen vented to drop the pressure. A 30 degree scope is passed inferior to the falciform ligament into the left upper quadrant over the lateral segment of liver. The post of the scope is positioned to the left, allowing visualization of the lesser curvature of the stomach with the end of the scope near the left aspect of the falciform.

A site for placement of the gastrostomy is selected on the skin, using a 22Ga needle as a finder. The site is marked, then infiltrated with local anesthetic and a 5mm transverse incision made.

Endoscopy is performed and the following noted:

Level in cm from the incisors, of the most proximal area of Barrett's esophagus.  
Level in cm from the incisors of the GE junction  
Appearance of the GE junction on retroflexed view. Extent of invasion of the tumor into the cardia or fundus.

A bite block is positioned (unless the patient is edentulous). The endoscope (usually a neonatal scope) is introduced into the esophagus and the video capture started. If the tumor will not allow passage of the scope, do not force the scope.

Once the scope is passed into the stomach, the fundus and duodenal bulb are suctioned.

Insufflation is then reduced to 4mm and the stomach insufflated to find the optimal location for tube placement which will minimize tension and will avoid injury to the right gastroepiploic artery. The reduced laparoscopic insufflation pressure allows endoscopic insufflation of the stomach.

Gastrostomy tube placement is done either by Pull or Seldinger technique.

### **Seldinger Gastrostomy**

Four T-fasteners are then used to affix the stomach to the anterior abdominal wall. These are arranged at 8:00, 10:00, 2:00, 4:00 relative to the proposed tube site. T-fasteners are not placed inferior to the tube site to avoid injury to right gastroepiploic vessels.

The J wire is then passed into the stomach, followed by the dilators, up to 20Fr. A 16Fr MIC gastrostomy tube is introduced and the balloon inflated with 5mL of *water*.

### **Pull Gastrostomy**

A 20Fr PULL PEG tube kit is opened. The snare and tube are passed to the upper operator. The snare is passed through the scope and opened in anticipation of the passage of the wire.

The angiocath from the kit is placed into the stomach through the abdominal wall. Once the snare has grasped the angiocath, the needle is withdrawn and the split wire passed through the Angiocath into the stomach. The snare is adjusted to grasp the split wire, which is pulled out through the mouth.

The laparoscopic port site is considered clean. The PEG tube, once it is pulled through the mouth, is considered dirty. The right abdomen is covered with a towel to protect the laparoscopic site.

The recording is now stopped. The split wire is joined to the PEG tube, which is pulled into place by the abdominal operator. The tapered portion of the tube will be the first source of resistance, which may require firm traction. The split wire (and PEG tube) is dropped of the table to the patient's left.

Once the tapered portion of the tube is through the abdominal wall skin, the next point of resistance will be the bumper of the PEG tube passing through

the tumor. In general, if a 5mm neonatal scope can pass the tumor, a 20Fr PEG tube can pass as well.

The tube is pulled into position and the measurement at the skin noted. The stomach is aspirated by the upper operator and insufflation is resumed at 8mmHg. Tension on the PEG tube is adjusted to allow apposition of the gastric serosa to the abdominal wall. If the tube is not easily apposed to the abdominal wall, T fasteners ("GI Anchors") must be employed.

The scope is withdrawn and the hypopharynx suctioned. The liquid from the 'First Step' disinfectant is suctioned through the scope, followed by water.

The abdomen is desufflated and the port sites closed with 4-0 Monocryl.





## Chapter 20

# Esophagectomy 1 Stage

### 20.1 Indications

MI Ivor Lewis (“One Stage”) esophagectomy is the most common approach to esophagectomy. The patient is positioned in ‘corkscrew’ position to allow simultaneous access to the abdomen and chest by prepping both abdomen and chest.

### 20.2 Room Prep

Confirm positioning equipment (Salo positioner ‘bucket’):

- Four black side-rail clamps
- Four rectangular lateral positioners
- Yassargil socket OR large-bore Clark socket
- Well-leg holder (used as an arm holder)

Critical supplies (check prior to pt in room)

- Stapler 25mm DSTXL
- Orvil 25mm
- Stapler 21mm DSTXL
- Orvil 21mm
- Echelon 60mm stapler with 10 gold loads and 2 gray loads
- Gel Port

Review initial PET images

#### **Anesthesia**

- Dual-lumen ETT tube (taped to left)
- Arterial line (left arm will be on arm board)
- Anesthesia may place paraspinal muscle blocks on right side.

### ##Position

Supine on blue foam pad.

Mark upper midline with non-sterile skin marker

Foley catheter (with Criticor temperature sensor)

Bovie pad

Lower-body Bair Hugger at level of thighs

Hair clipped from abdomen, right chest, and right axilla.

Pre-existing jejunostomy

- Prep into field
- Remove any eschar at site
- Secure to skin inferiorly with 0 silk

Shoulders are shifted to the right in preparation for ‘corkscrew’ positioning

Lateral positioners positioned with pad extending from greater trochanter inferiorly.

Velcro strap over thighs and over lateral positioners

Yassargil socket attached to headpiece of bed on left side.

Well-leg holder attached to Yassargil socket and used to support right forearm (which will cross body). Right arm crosses body and is supported on well-leg holder.

Lateral positioner placed posterior to spine and scapula to support right chest, allowing access for thoracoscopy.

Arm holder is dropped towards the floor enough that right arm is brought forward.

### Prep

Chloroprep (two sticks) of abdomen, right chest, right axilla. Particular attention to prepping as far as possible to the left lateral side and the right lateral side. Nipple prepped into field.

### Drape

Proximal right arm draped with 1000 (clear adhesive) drape.

Two blue plastic “U” drapes with center of the “U” on either lateral side with tails forming the perimeter of the field Trauma drape. All of right chest and

axilla is kept within the field, as is the lateral aspect of the left upper quadrant. The field does not need to extend inferior to the umbilicus. In general, it is usually possible to keep all of these area in the field without cutting the drape, except in very large patients. Ioban strips (4") around the periphery of the surgical field after the trauma drape. Laparoscopic cords (gas, light cord, camera) to tower at left shoulder. Suction irrigator brought off field. Laparoscopic LigaSure (2 bars) and Bovie (30/30).

### 20.2.1 Time Out

Operation header on consent

Tumor location and likelihood of division of esophagus from abdomen (two-phase operation) vs division of esophagus from chest (four-phage operation).

Blood:

- Surgeon's expectation of blood loss
- Availability of blood (type/screen vs type/cross).

*Comorbidities:*

Cardiopulmonary disease. If echo, ejection fraction and aortic valve area (if abnormal)

Beta blockage: Note whether patient on home beta blockade and if so, whether home medication was taken the morning of surgery.

Anticipated Intraop Problems:

- Possibility of tension left pneumothorax due to carbon dioxide entry into left chest during mediastinal dissection
- Expectation of ventilatory difficulties due to carbon dioxide entry into the right chest Gastric mobilization – Greater Curvature

## 20.3 Gastric Mobilization

After infiltration of 1% marcaine, 7cm upper midline incision made for handport. Small incision is made and pylorus palpated, if possible. Incision is centered over pylorus. GelPort inserted, and abdomen insufflated to 15mmHg. Two 5mm ports placed LUQ. Medial LUQ 5mm port placed in angle between left costal margin and superior edge of GelPort ring. Lateral LUQ 5mm port placed as far lateral as possible. Depending upon visualization, a third port may be required between these two and somewhat more inferior.

If feasible, division of gastrocolic omentum starts by delivering transverse colon into GelPort and dividing ligament with cautery and LigaSure in the avascular plane just cephalad to transverse colon. Dissection proceeds as far proximal and distal and feasible. It is important to avoid damage to the right gastroepiploic artery.

Colon is returned to abdomen and gastrosplenic ligament divided going using LigaSure, taking care to avoid the colon and the right gastroepiploic artery. For patients with a bulky omentum, it may be helpful to place an additional port in the left mid-quadrant for the camera to facilitate dissection of omentum off the transverse colon. Stomach is retracted to the patient's right with the back of the left hand, placing the gastrocolic ligament (and short gastric arteries) on stretch. Left gastroepiploic artery divided with LigaSure near its origin. Short gastric arteries divided with Ligasure close to spleen. As superior aspect of short gastrics is reached, the dissection plane shifts medially to create a tunnel towards the base of the left crus. This places the most superior short gastric vessels on stretch and facilitates their division. Once all short gastric vessels divided, peritoneum tethering fundus to diaphragm is incised, and fundus brushed medially.

### 20.3.1 Distal mobilization

Attention is directed to mobilizing the lateral aspect of the duodenum. This can either be accomplished with the camera through a LUQ port and a 45 degree camera or by placing an additional 5mm port in the RLQ. Hook cautery is used to incise the connective tissue lateral and posterior to the duodenum. The gastrocolic ligament is now dissected distally, taking care to preserve the integrity of the right gastroepiploic vessels. An areolar plane generally exists between the fat pad containing the right gastroepiploic vessels and that containing the transverse mesocolon vessels.

### 20.3.2 Left gastric artery

Lymph nodes around the celiac axis and left gastric artery are now dissected. The extent of dissection depends upon the tumor location and the presence of nodes here either based upon imaging or palpation. Dissection begins on the superior edge of the pancreas, and proceeds superiorly to the right crus. The left gastric (coronary) vein is usually located to the left of the artery and is divided with the LigaSure. In a two-phase approach, the left gastric artery is now divided with a 30mm gray load (2.0mm) Echelon linear stapler. For patients with mid-esophageal tumors, (for whom a four-phase approach is used), if there is any question about the resectability of the tumor, division of the left gastric artery is generally deferred until the second abdominal phase.

### 20.3.3 Mediastinal dissection

The esophagus is now dissected circumferentially at the gastroesophageal junction. The peritoneum overlying the diaphragm is incised, and circumferential dissection of esophagus performed. On the left side, it is helpful to distract the left crus laterally with a Prestige clamp placed on the left crus. The right pleura is widely entered in order to both facilitate the thoracic dissection of the esophagus and placing the conduit into the right chest in preparation for the final thoracic phase. Division of Esophagus (Two-Phase only) In a four-phase approach, the esophagus is divided from the right chest during the first (of two) thoracic phases. In a two-phase approach, the esophagus is divided from the abdomen by reaching up into the mediastinum to divide the esophagus above the tumor. This is only feasible for tumors of the gastroesophageal junction. In a two-phase approach, the esophagus is divided with a 60mm Medium-Thick (Gold) Echelon TriStaple stapler.

### 20.3.4 Division of Esophagus (Four-Phase)

Penrose Drain (Four-Phase only) In a four-phase approach, the esophagus is divided from the chest. In order to facilitate the thoracic dissection, a 1/4 penrose drain is tied around the distal esophagus and the drain is slid cephalad into the mediastinum. The 'tails' of the drain are directed into the right chest so that they can be grasped from the right chest during the thoracic phase and can be used to provide traction on the esophagus.

### 20.3.5 Entry into right chest

The pneumoperitoneum in the abdomen is vented and the gas pressure turned down to 8mmHg in preparation for the thoracic phase. The bed is rotated to the left 20 degrees. A site for entry into the left chest is selected just posterior to the tip of the scapula. An incision is made here and a 5mm optical port with a 5mm 0 degree scope used to enter the chest. The chest is insufflated at 8mmHg of carbon dioxide, which helps to both collapse the lung and depress the diaphragm. Two 12mm ports are placed. The more superior is placed lateral and superior to the nipple. The inferior port is placed just lateral to the diaphragmatic reflection. It is critical to avoid injury to the diaphragm (and liver) with the inferior port placement. A mini-thoracotomy incision is placed along the mid-axillary line, frequently in the same interspace as the inferior/anterior 12mm port. The chest is entered just superior to the rib and the intercostal muscles divided with the LigaSure device to allow the ribs to separate. A narrow Deaver retractor is used to gauge the space between the ribs, as the width of the retractor approximates the diameter of the 25mm stapler. A 5mm 'U' port is generally placed as high as possible midway between the scapular tip port and the anterior/superior 12mm port to the 4 right 5

**Division of Esophagus (Two-Phase)** In patients with low-lying tumors, it is possible to divide the esophagus above the tumor from the abdominal approach. This evaluation is facilitated by review of the preoperative endoscopy (and EUS), and the PET scan, particularly the PET obtained prior to neoadjuvant chemoradiation.

After dissection of the mediastinum, a Echelon stapler with a 60mm Medium-Thick (Gold) load is inserted through a 12mm port either placed either through the GelPort or by upsizing the most lateral LUQ

**Construction of Conduit** The distal esophagus and stomach is exteriorized through the GelPort. The lesser curvature vessels are divided with the LigaSure 7-9cm cephalad from the pylorus. The stomach is now placed on stretch along the greater curvature. An Echelon Medium-Thick (Gold) stapler is used to construct a 5-6cm wide gastric tube. In constructing the conduit in a patient with a tumor of the GE junction invading into the cardia, it is important to be certain that the staple line to construct the conduit stays clear of the tumor. Patients with tumors invading the cardia are at risk of a positive distal margin, meaning that microscopic tumor may be left in the wall of the gastric conduit. To make things more complicated, in patients with low-lying esophageal or GE junction tumors, not all of the length of the conduit are needed in order to reach to the level of the esophageal transection. After construction of the anastomosis, the 'extra' cephalad portion of the conduit (near the angle of His) are excised as the 'additional gastric margin.' In order to distinguish which portions of the conduit staple line which will be used to replace the esophagus and those which will be included in the 'additional gastric margin', both sides of the gastric conduit staple line are marked with sutures designated 'A', 'B', 'C', etc proceeding from the Angle of His to the antrum.

The distal esophagus and GE junction are now sent for frozen section.

**Feeding jejunostomy** The ligament of Trietz is identified and the jejunum is identified 20cm distal and marked with a directional suture. A site is selected to the left of the handport incision. A 16Fr Cook Introducer Kit is used to pass a 16Ga needle, followed by a J wire, through the left rectus muscle. A skin incision 4mm in length is made adjacent to the J wire. The 16Fr dilator and sheath are now passed through the rectus muscle and the dilator and wire removed. A 14Fr Jejunostomy Tube 0301-14 is selected and the 'wings' trimmed off with a scalpel. The jejunostomy tube is passed through the peelaway sheath, which is removed.

A pursestring suture 1.5cm in diameter is placed on the antimesenteric border of the jejunum. The pursestring is started and ends on the lateral aspect. A second 16Fr Cook Introduced Kit is used to introduce the J wire through the center of the pursestring.

**Placement of Drains** Two (or three) 19Fr full-fluted Blake drains (72230) are placed:

JP1: placed into the left pleura through the hiatus. Drain is brought out through the most lateral 5mm port site on the LUQ JP2: placed into the right pleura through the hiatus. Drain is brought out through the next most medial 5mm LUQ port site JP3: (optional) placed in the abdomen posterior to the left lateral segment of the liver and brought out through the most medial 5mm LUQ port site Transposition of Conduit The gastric conduit is now placed into the right pleura through the mediastinum, with the assistance of a laparoscopic Babcock and gentle pressure on the greater curvature with the fingers.

Entrance into the R chest (Two Phase) For two-phase operations, ports are now placed into the right chest, as stated above . In similar fashion, the inferior pulmonary ligament is dissected and the lung reflected anterior.

Anastomosis The right chest is entered and the right lung reflected anterior with the paddle placed in the superior/anterior port. The gastric conduit is placed into the mediastinum by tucking it medially from the right pleura into the posterior mediastinum, in order to allow the conduit to take the most direct path from the hiatus to the proximal esophagus. Gentle superior tension is now applied to the conduit in order to eliminate redundancy. The paddle retractor is now moved from the anterior/inferior port to the anterior/superior 12mm port and the lung reflected anterior and inferior.

OrVil The OrVil device is now used to place a EEA anvil into the distal esophagus. In general, a 25mm size is selected, unless the patient has particularly small frame, in which case a 21mm size is used. Two stay sutures of 2-0 silk on RB-1 needles are placed in the center of the esophageal staple line 3mm apart. A Harmonic scalpel is used to divide the staple line. The OrVil device is passed through the mouth and is passed through the fenestration in the staple line. The anvil portion of the OrVil is oriented so that the rounded portion is placed against the roof of the mouth. The OrVil is guided into the hypopharynx by pulling on the the tube end of the device. As the anvil approaches the hypopharynx, the jaw is pulled forward to allow passage of the anvil.

The shaft of the anvil is brought through the esophageal staple line, and the tube disconnected from the anvil by cutting the blue sutures.

The superior end of the conduit is opened along the staple line and the DST XL stapler (matching the diameter of the OrVil) introduced through the Alexis device. The stapler shaft is placed into the open end of the gastric conduit and the conduit pulled over it ('sock over shoe'). The stapler spike is brought out through the greater curvature. The anvil is grasped with a Maryland grasper placed through a superior 5mm port and the two components of the stapler are mated and the stapler tightened and fired. The knob of the stapler is rotated two turns counter-clockwise until a click is felt, at which time the anvil will flip. The stapler is withdrawn and the donuts examined and sent for pathologic exam.

The anastomosis is completed by firing a Echelon Medium-Thick (Gold) linear across the conduit cephalad to the anastomosis. The excess conduit is sent for

pathologic exam as ‘additional gastric margin.’

NG Tube A Covidien Salem Sump 18Fr nasogastric tube is passed by the anesthetist. A laparoscopic BK ultrasound is used to monitor the passage of the NG tube through the esophagus and into the gastric conduit. The NG tube is passed to the level that all four dots are outside the nose, with the 4th dot at the nares. The NG tube is secured with an AMT bridle.

Chest Tube A 28Fr Blake chest tube is placed through the anterior/inferior 12mm port and is positioned into the posterior mediastinum. JP2 is placed near the gastric conduit. The right lung is re-inflated.

Closure The stapler access port incision is closed with 0 Vicryl to approximate the serratus muscle. The incisions are closed with 4-0 Monocryl followed by Dermabond.



# Esophageal Cancer



## Chapter 21

# Esophageal Overview

Esophageal cancers can be grouped into 4 treatment categories:

- Superficial → Endoscopic therapy
- Localized → Primary surgery
- Locally Advanced → Trimodality therapy
- Metastatic → Systemic therapy

Patients with minimal dysphagia, no weight loss, and small (<3cm length) tumors are evaluated with endoscopic ultrasound:

- If uT1 on EUS and <2cm in size, endoscopic mucosal resection yields more information and may be therapeutic for tumors with negative margins and without high-risk features.
- If uT2N0 on EUS, and PET scan shows a small tumor (MTV <10cm<sup>3</sup>), primary surgery is preferred in patients who are good surgical risks
- If T3 or N+ on EUS, if PET shows no metastatic disease, trimodality therapy is optimal)

Patients with dysphagia to solids or weight loss or tumor length >3cm are unlikely to have T1-2 tumors and can be evaluated with PET scan.

- If PET shows disease confined to the esophagus and regional nodes, trimodality therapy (chemoradiation followed by surgery) is optimal.
- If PET shows metastatic disease, patients are eligible for palliative chemotherapy with radiation for treatment of symptoms of dysphagia.
- If PET shows extra-regional lymph node disease, patient is at high risk for distant disease and can be treated with induction chemotherapy followed by chemoradiation and surgical evaluation.



## Chapter 22

# Staging

The staging workup begins once a diagnosis is made on endoscopy.

The first step is to make a preliminary determination whether the tumor is early stage (and can be treated with endoscopy or primary surgery) or later stage (and treated with chemoradiation followed by surgery or with)

The diagnostic studies needed for these treatment groups are different, so the workup can be made more efficient by sorting patients at presentation in to two groups:

Patients with minimal dysphagia, no weight loss, and tumors with less than 3cm cranio-caudal extent have a reasonable chance of being T1 or T2 tumors. Tumors <3cm in length are much more likely to represent T1-2 lesions than those  $\geq 3$ cm (Hollis et al., 2017)

Superficial and Localized tumors generally present with minimal dysphagia or weight loss. These tumors may present with bleeding, or dysphagia without weight loss. For these patients, determining the precise T stage is important in their workup, so **endoscopic ultrasound** is the most frequent staging study after diagnosis.

Locally-advanced or metastatic tumors tend to present with dysphagia and weight loss. At first approximation, these tumors are usually clinical T3 lesions, and the important bifurcation in their treatment is the presence or absence of metastatic disease. For patients with dysphagia and weight loss, **PET** is the most frequent initial staging study after diagnosis.

Patients who present with dysphagia are likely to have T3 or T4 disease, which is generally treated with neoadjuvant chemoradiation followed by surgery. Data from Memorial Sloan Kettering [Ripley 226] among 61 patients with esophageal cancer who presented with dysphagia, 54 (89%) were found on EUS to have uT3-4 tumors. On the other hand, among 53 patients without dysphagia, 25 (47%) were uT1-2, and were potentially candidates for primary surgery. Their

conclusion was that EUS could be omitted from the workup of patients with dysphagia, but is useful in patients without dysphagia.

PET can be helpful in evaluating patients who may have T1-2 disease, and might be candidates for primary surgical therapy. A comparison of PET and EUS [malik,claxton,1] showed that uT1-2 tumors had median metabolic tumor volume (MTV) of  $6.7\text{cm}^3$ , compared with uT3-4 tumors, with a median SUV of  $35.7\text{cm}^3$ .

## Chapter 23

# Superficial EsoCa

Superficial esophageal cancer is usually asymptomatic, which means that the diagnosis is generally made in the context of surveillance for Barrett's esophagus.

Nodular Barrett's esophagus can be best evaluation with endoscopic mucosal resection, which can provide further staging information if an adenocarcinoma is found, such as depth of invasion, differentiation, and lymphovascular invasion.

Larger lesions should first be evaluated with endoscopic ultrasound (EUS)?

EUS is less sensitive for T1 lesions (Bergeron et al., 2014) -> use EMR for diagnosis (Maish and DeMeester, 2004)

(Should nodular Barrett's be evaluated with EUS prior to EMR?)

T1a tumors have a low risk of nodal metastasis (Dunbar and Spechler, 2012)

### 23.1 Endoscopic Mucosal Resection (EMR)

For patients with nodular Barrett's esophagus or small tumors judged to be T1 by endoscopic ultrasound, endoscopic mucosal resection (EMR) can be diagnostic and potentially curative.(Thomas et al., 2009)

EMR also helps establish the difference between T1a and T1b compared with pathology (Worrell et al., 2018)

EMR is likely sufficient for small tumors with favorable pathologic factors(Pech et al., 2014) (Nurkin et al., 2014):

- Size less than 2cm
- Lateral and deep margins clear
- Absence of lymphovascular invasion

- Well- or moderately- differentiated

EMR: (Soetikno et al., 2005)

See Molina JTCVS 153:1206

EMR for high-grade dysplasia (Shaheen et al., 2009)

EMR for low-grade dysplasia (Phoa et al., 2014) resulted in 25% risk reduction in progression to HGD.

Endoscopic submucosal dissection is a technique for deeper endoscopic removal of esophageal lesions using endoscopic cautery, which dissects through the submucosa. ESD has a higher rate of curative resection (Cao et al., 2009) albeit at the cost of prolonged operative times and increased risk of complications such as bleeding. (Repici et al., 2010)

ESD takes more time and has higher R0 resection rate but similar recurrence rate at 2 years (Terheggen et al., 2017)

Need for RFA of Barrett's after EMR: (Haidry et al., 2013) Combination therapy with EMR and RFA results in lower rate of recurrence than EMR alone. (Pech et al., 2008)

RFA for Barrett's national registry (Ganz et al., 2008)



## Chapter 24

# Localized EsoCa

### 24.1 T1b Tumors

### 24.2 T2N0 Tumors

Multiple studies have failed to show the additional benefit of chemotherapy or chemoradiation for pT2N0M0 esophageal cancer patients treated with radiation.

Neoadjuvant chemo not likely to be helpful for early stage disease - FFCD 9901 [Marette 2416] enrolled patients with T1-2 or T3N0 tumors to chemoradiation followed by surgery versus surgery alone. The majority of the tumors (72%) were squamous cell carcinoma. Postoperative mortality was significantly increased in the chemoradiation arm (11.1% vs 3.4%).

Meta-analysis of 5265 patients in 10 studies showed that while neoadjuvant therapy was associated with a reduction in positive margin rate, there was no difference in terms of recurrence or survival. [Mota 176]

French trial FREGAT (Markar et al., 2016)

Retrospective review of the National Cancer DataBase failed to demonstrate a difference in survival of cT2N0M0 esophageal cancer with or without preoperative chemoradiation. (Speicher et al., 2014)

A retrospective report from Johns Hopkins examined outcomes of T2N0 squamous cell carcinoma patients and showed equivalent outcomes for primary surgery vs neoadjuvant chemoradiation followed by surgery (Zhang et al., 2012)

## 24.3 Staging of T2N0 Tumors

The challenge for treatment decision-making is the limited sensitivity of endoscopic ultrasound in ruling out pT3 or pN+ disease. In other words, if a patient who is thought to have cT2N0 disease undergoes resection, and is found on pathology to have pT3 or pN+ disease, this would dictate the need for post-operative chemoradiation. In general, chemoradiation after esophagectomy is difficult for patients to tolerate, with a \_\_\_\_ % chance of failure to complete therapy.

Data from the Cleveland Clinic looked at 53 patients judged to be T2N0 by endoscopic ultrasound (uT2N0) were treated with primary surgery. Pathologic examination showed that 17 (37%) were understaged by endoscopic ultrasound, and were pathologic (pT3) in 4 or node positive (pN+) in 13 cases. These patients were treated with postoperative adjuvant chemoradiation. (Rice et al., 2007)

It is critical, therefore, in patients for whom primary surgery is contemplated, to attempt to identify those with occult T3 or N+ disease.

Patients who appear to have limited stage disease benefit from evaluation with a combination of

See also PMID:25047477

(MTV)

(Tumor Length)

(dysphagia)

##Primary Surgery {#primary\_surgery}

NCCN recommends PET scanS

Most common sites of metastasis are liver, lung, bones, adrenal.

PET detects occult metastasis in 10-20% of cases (Kato et al., 2002, Kim et al. (Apr)). Among 129 patients with esophageal cancer, PET detected additional sites of disease in 41% and changed management in 38% (Chatterton et al., 2009)

PET for restaging detects interval development of metastatic disease in 8-17% of cases (van Vliet et al., 2008)

## Chapter 25

# Locally Advanced EsoCa

Tumors that are T2N<sup>+</sup>M0 or T3NxM0 are considered locally-advanced. The high rate of failure with surgery alone has led to development of adjunctive therapies.

### 25.1 Trimodality Therapy

Trimodality therapy consists of chemoradiation followed by surgery.

CROSS trial randomized 364 patients with resectable esophageal and gastroesophageal junction tumors (75% adenocarcinoma) to neoadjuvant chemoradiation consisting of 4,140 cGy of radiation with concurrent carboplatin and paclitaxel or surgery alone.(van Hagen et al., 2012) Clinical node-positive disease was present in 16%. Pathologic complete response was seen in 23% of adenocarcinoma and 49% of squamous cell carcinomas. Median overall survival was 49 months after trimodality vs 24 months after surgery alone (p=0.003). Squamous cell carcinomas appeared to have particular benefit, with a hazard ratio of 0.42 for squamous cell vs 0.74 for adenocarcinoma. Median survival was improved for adenocarcinoma from 27.1 months to 43.2 months, but the median survival for squamous cell increased from 27.1months to 81.6 months for squamous cell. Rate of R0 resection was higher with chemoradiation (92% vs 69% p<0.001) and local recurrence rates lower (14% vs 34% P<0.001), and peritoneal recurrence lower (4% vs 14% P<0.001). Despite the relatively low dose of radiation, in-field recurrences were less than 5%. The primary cause of failure was distant disease (31%) and local/regional failure (14%).(Oppedijk et al., 2014)

Alternative to carboplatin is FOLFOX (SOG trial (Leichman et al., 2011))

Ongoing PROTECT trial compares FOLFOX to paclitaxel and carboplatin (Messager et al., 2016)

See also Definitive ChemoRT

### 25.1.1 Neoadjuvant chemoRT for SCCA

NeoCRTEC5010 (Yang et al., 2018)

Meta-abalysis of chemoRT vs chemo (Zhao et al., 2018)

### 25.1.2 Neoadjuvant chemotherapy followed by surgery

POET Trial (Pre-Operative therapy in Esophageal adenocarcinoma Trial) treated patients with adenocarcinoma of the gastroesophageal junction with either neoadjuvant chemotherapy (5-FU, leucovorin, cisplatin) followed by surgery or induction chemotherapy with the same agents, followed by chemoradiation (4000cGy with concurrent cisplatin and etoposide). The study failed to meet its accrual goal, but there was a suggestion of improved 3-year survival with preoperative chemoradiation (47.4% vs 27.7%  $p=0.07$ ) as well as improved local control (76.5% vs 59%). In addition, chemoradiation was associated with a higher pathologic complete response rate (15.6% vs 2%)(Stahl et al., 2009). A meta-analysis of 33 randomized trials further suggested a greater benefit from neoadjuvant chemoradiation followed by surgery compared with neoadjuvant chemotherapy followed by surgery(Pasquali et al., 2017) and a similar meta-analysis (Sjoquist et al., 2011)

#Active Surveillance

EGD is poor predictor of pCR (Sarkaria et al., 2009)

## 25.2 GE Junction

(Siewert et al., 2006)

## 25.3 Induction chemotherapy followed by chemoRT

See NCCN pages M-25 and M-26

Stahl (Stahl et al., 2009) randomized patients to preoperative chemotherapy (A) vs preoperative chemotherapy followed by preoperative chemoradiation (B). Higher pCR rate in arm B (15.6% vs 2%) and ypN0 resection (64.4% vs 37.7%).

## 25.4 Postoperative chemoradiation

Intergroup-0116 (Macdonald et al., 2001) (Smalley et al., 2012) treated 556 patients with adenocarcinoma of the stomach or GE junction with surgery alone vs surgery followed by postoperative chemoradiation. After a median followup of over 5 years, median overall survival in the surgery alone group was 27 months vs 36 months in the postoperative chemoradiation group ( $p=0.005$ ) Decrease in local failure as the first site of failure in the chemoradiation group (19% versus 29%).

Chemoradiation after resection of GE junction tumors (Kofoed et al., 2012) among a group of 211 patients with GE junction adenocarcinoma with positive lymph nodes with improved 3-year disease-free survival (37% vs 24%).



## Chapter 26

# Chemoradiation

**\*\***This section addresses chemoradiation as primary (definitive) therapy for esophageal cancer. See also Trimodality Therapy

### 26.1 Phase II Studies

Experience with patients who refuse surgery or are medically unfit:

MD Anderson report of 61 patients out of 622 tri-modality-eligible patients who refused surgery after cCR. 5-year overall survival was 58%. 13 developed local recurrence during surveillance, and 12 had successful salvage esophagectomy (Taketa et al., 2012). The same group compared those who underwent surgery vs definitive chemoradiation in a propensity-matched fashion (Taketa et al., 2013). Irish study of 56 patients aged 70 or older treated with neoadjuvant chemoradiation followed by selective surgery. Median survival was 28 months overall, 47 months for those with cCR, 61 months for primary resection, 46 months for cCRs who did not undergo resection, and 29 months for those with salvage esophagectomy (Furlong et al., 2013).

Castoro (Castoro et al., 2013)

preSANO (Chirieac et al., 2005) Clinical Response evaluation after chemoRT for esophageal cancer with PET and EGD.

### 26.2 ChemoRT vs Trimodality therapy

The sensitivity of squamous cell carcinoma of the esophagus to chemoradiation has raised the question whether

Stahl Locally advanced squamous cell carcinoma randomized to induction chemotherapy (cisplatin, etoposide, 5FU with leucovorin) followed by chemoradiation (4000cGy with concurrent cisplatin and etoposide) followed by surgery compared with induction chemotherapy followed by chemoradiation (6400cGy with concurrent cisplatin and etoposide).(Stahl et al., 2005) progression-free survival was better in the trimodality group (64.3% vs 40.7%) Treatment-related mortality was substantial in the surgery arm (13% vs 4%). This would be considered an excessive rate of operative mortality by modern standards. Unsurprisingly, there was no difference in overall survival between groups, in part because the surgical group had an excess 9% mortality rate from treatment. Two-year survival in the surgery arm was 40% vs 35% in the definitive chemoradiation arm.

In the French FFCO trial, 444 patients with carcinoma of the esophagus (90% squamous cell) were treated with two cycles of 5-FU and cisplatin with concurrent radiation.(Bedenne et al., 2007) Patients with a partial or complete clinical response to chemoradiation were randomized to either surgery or a boost of radiation. Patients who did not respond to chemoradiation were treated with surgery and were eliminated from the study. Only 259 of the original 444 patients (59%) went on to randomization, with the remainder (those not responding to chemoradiation) treated with surgery. Of the randomized group, median survival was 17.7months in the surgery arm versus 19.3months in the definitive chemoradiation arm. Like the Stahl study, treatment-related mortality in the surgical arm was high (9% versus 1%).

#Active Surveillance

EGD is poor predictor of pCR (Sarkaria et al., 2009)



## Chapter 27

# Radiation

RTOG 94-05 clinical trial (Minsky et al., 2002)



## Chapter 28

# Esophagectomy

Three general approaches exist for surgical therapy.

Trans-thoracic or Ivor Lewis esophagectomy(Visbal et al., 2001) removes the intrathoracic portion of the esophagus and constructs an anastomosis within the chest. The approach include an abdominal phase, during which an esophageal substitute is constructed (usually from stomach). A thoracic phase then removes the intrathoracic esophagus and constructs an anastomosis within the chest cavity.

A McKeown esophagectomy utilizes three surgical fields: abdomen, right chest, and neck. The right chest approach allows dissection of peri-esophageal lymph nodes, and the cervical incision allows removal of the total esophagus.(McKeown, 1976) This approach is useful for tumors which involve the proximal thoracic esophagus, to ensure a negative margin. The cervical anastomosis carries a higher risk of anastomotic leak than a thoracic anastomosis, although the morbidity of a cervical anastomosis leak is less serious than that of a leak of a thoracic anastomosis.

A transhiatal esophagectomy approaches the esophagus from the abdomen through the hiatus and from neck. By blunt dissection the esophagus is freed up without the need for thoracotomy. An esophageal substitute is then brought from the abdomen to the neck through the mediastinum(Orringer and Sloan, 1978) (Orringer et al., 2007) – Orringer Ann Surg 2007 –> The operation is designed to avoid the pulmonary toxicity of the right chest approach. On the other hand, the blunt nature of the mediastinal dissection means that fewer lymph nodes are harvested than with a trans-thoracic approach.

Randomized trial of transthoracic esophagectomy with extended lymph node dissection versus transhiatal esophagectomy showed fewer pulmonary complications with the transhiatal approach. (Hulscher et al., 2002) Fewer lymph nodes were harvested with a transhiatal approach. A post-hoc analysis showed that

among patients with 1-8 positive lymph nodes, survival with improved with the extended lymph node dissection.(Omloo et al., 2007)

Minimally-invasive approaches to esophagectomy are now common, with evidence for less perioperative morbidity than an open approach (Biere et al., 2012) (Zhou et al., 2015)

Randomized trial of a hybrid MIE (with laparoscopy and thoracotomy) was associated with lower postoperative complications than open esophagectomy (Mariatte et al., 2019)

High volume (Birkmeyer et al., 2003) (Wouters et al., 2009)

Siewert III lesions are considered gastric cancers (Rusch, 2004) (Siewert et al., 2006)

Laparoscopy may be helpful in Siewert III tumors (de Graaf et al., 2007)

### 28.0.1 Preoperative Evaluation

Dysphagia can be scored according to Mellow et al (Mellow and Pinkas, 1985):

- 0 No dysphagia
- 1 Dysphagia to normal solids
- 2 Dysphagia to soft solids (ground beef, poultry,fish)
- 3 Dysphagia to solids and liquids
- 4 Inability to swallow saliva

## 28.1 Minimally-invasive Esophagectomy

Higher lymph node yield with MIE vs open approach (Kalff et al., 2020)

## **28.2 Transthoracic**

## **28.3 Transhiatal**

## **28.4 Three-hole**

## **28.5 Extended lymphadenectomy**

## **28.6 Early Recovery Pathways**

ERAS Society Guidelines

## **28.7 Salvage esophagectomy**

(Markar et al., 2014)

(Swisher et al., 2002)



## Chapter 29

# Metastatic EsoCa

### 29.1 Palliative radiation

Palliative radiation vs chemoradiation (Penniment et al., 2018)

Radiation alone favored over chemoradiation in the palliative setting (Penniment et al., 2018)

### 29.2 Chemoradiation vs chemotherapy in Stage IV

(Guttmann et al., 2017)

### 29.3 Stents for malignant disease

(Vakil et al., 2001)

Review of guidelines 2010 Am Society GI (Sharma et al., 2010)





## Chapter 30

# Survivorship

### 30.1 Nutritional consequences of esophagectomy

#### 30.1.1 Vitamin D deficiency

Vitamin D deficiency is defined as serum 25(OH)D levels below 20ng/mL

- Replacement with 2000 IU Vitamin D3 daily (Khan and Fabian, 2010)

Vitamin D insufficiency is defined as serum 25(OH)D level 20-29ng/mL

- Replacement with 1000-2000 IU Vitamin D3 daily

NIH ODS Vitamin D info

(Baker et al., 2016)

Weight loss (Martin and Lagergren, 2009) (Ouattara et al., 2012)

### 30.2 Cardiac toxicity of radiation

(Beukema et al., 2015) (Frandsen et al., 2015) (Gharzai et al., 2016)

### 30.3 Surveillance

**T1a treated with endoscopic resection** EGD every 3 mo for first year, then every 6 months for second year, then annually (Shaheen et al., 2016)

**T1b treated with endoscopic resection** EGD every 3 mon for first year, then every 4-6 months for second year, then annually CT chest/abdomen every 12 months for up to 3 years (as clinically indicated)

**T1b treated with esophagectomy** EGD every 3-6 months for first 2 years, then annually for 3 more years. CT every 6-9 months for first 2 years, then annually up to 5 years.

**Stage II or III treated with chemoradiation.**

These patients are at risk for local recurrence (Sudo et al., 2014) and some may be candidates for salvage esophagectomy. Most relapses (95%) occur within 24 months. See also (Taketa et al., 2014)

**Locally-advanced treated with trimodality therapy**

Local/regional relapses are uncommon. (Dorth et al., 2014) (Oppedijk et al., 2014) (Sudo et al., 2013) => NCCN does not recommend EGD. 90% of relapses occur within 36 months of surgery.

CT every 6 months up to 2 years (if patient is a candidate for additional curative-intent therapy)

# Gastric Cancer



## **Chapter 31**

# **Gastric Overview**



## Chapter 32

# Superficial Gastric





## Chapter 33

# Locally-Advanced Gastric

Locally-advanced gastric cancer (T3 or N<sup>+</sup>) is generally treated with some form of adjuvant therapy, which has been shown to improve upon the outcomes with surgery alone.

### 33.1 Preoperative Chemotherapy

FLOT chemotherapy (Al-Batran et al., 2019)

MAGIC study randomized 503 patients to perioperative ‘sandwich’ therapy consisting of epirubicin, cisplatin, and 5-FU versus surgery alone. In the perioperative chemotherapy group, 4 cycles were administered prior to surgery, and 4 cycles afterwards. Tumors of the esophagus or gastroesophageal junction comprised 26% of the study population. While over 90% of patients assigned to the chemotherapy arm completed their preoperative chemotherapy, only 66% completed their postoperative therapy. Survival at 5 years was 36% in the perioperative chemotherapy group, compared with 24% in the surgery group ( $p < 0.001$ ). (Cunningham et al., 2006)

CLASSIC clinical trial randomized 1033 patients with stage II or III gastric cancer after D2 gastrectomy to 6 months of adjuvant chemotherapy versus surgery alone. Three-year survival was improved in the chemotherapy group (74% *v* 59%). (Bang et al., 2012)

The FFCO trial randomized patients to preoperative chemotherapy with 2 or 3 cycles of cisplatin and 5-FU versus surgery alone. Tumors of the lower esophagus or gastroesophageal junction comprised 75% of the study population. Survival at 5 years was longer in the chemotherapy group (38%) versus 24% in the surgery alone group ( $p = 0.02$ ). (Ychou et al., 2011)

### 33.2 Postoperative chemotherapy

CLAASIC trial (Noh et al., 2014) (Bang et al., 2012) patients with II or IIIB gastric cancer received gastrectomy with D2 node dissection randomized to postoperative chemotherapy with capecitabine and oxaliplatin. Chemotherapy group had improved 3-year DFS (74% vs 59%  $P < .0001$ )

### 33.3 Postoperative chemoradiation

Intergroup 0116 trial (Macdonald et al., 2001) Surgical quality control was poor, as 90% were treated a limited lymph node dissection. Long-term followup, however (Smalley et al., 2012) showed a persistent benefit of postoperative chemoradiation.

ARTIST trial 450 patients treated with a D1  $\alpha$  gastrectomy were randomized to adjuvant capecitabine and cisplatin versus chemoradiation consisting of two cycles of capecitabine/oxaliplatin followed by chemoradiation followed by chemotherapy. Overall 3- year survival did differ between groups (78.2% vs 74.2%  $p = 0.86$ ). A post-hoc analysis of patients with positive nodes showed a beneficial effect of chemoradiation (77.5% v 72.3%  $p = 0.365$ ). (Lee et al., 2012)

CRITICS trial treated all patients with preoperative chemotherapy followed by surgery. Postoperative patients were then randomized between additional chemotherapy versus chemoradiation.

### 33.4 Preoperative chemoradiation

(Ajani et al., 2006)

## Chapter 34

# Hereditary Diffuse Gastric Cancer



# Colon Cancer



## Chapter 35

# Colorectal Overview

### Resources

ASCRS Guidelines for treatment of Colorectal Cancer 2022

ASCRS Guidelines for Early Recovery





## Chapter 36

# Colon Cancer Staging

### 36.1 CEA level

CEA should be routinely checked prior to surgery

### 36.2 CT Scan

### 36.3 PET scan

PET is not recommended routinely for evaluation of colon cancer.(Vogel et al., 2022)

#### 36.3.1 Indications for PET in colorectal cancer:

- Workup of rising CEA
- Indeterminate extra-hepatic findings on CT or MRI
- Evaluation of possible local recurrence

### 36.4 Adjuvant Therapy

Adjuvant therapy for stage II colon cancer Review {28399381}



## Chapter 37

# Stage IV Colon Cancer

### 37.1 Carcinomatosis

### 37.2 Colon resection in stage IV colon cancer

MSKCC: Colon surgery patients with bilobar liver metastasis treated with chemotherapy with primary tumor in place. Surgical intervention needed in 7% of cases (Poultides and Paty, 2011)

### 37.3 Immunotherapy for dMMR (MSI high)

Colorectal cancers deficient in mismatch-repair proteins (dMMR) respond poorly to 5FU-based chemotherapy but do respond to immune-checkpoint inhibitors ( $\alpha$ -PD-1, ( $\alpha$ -PD-L1, ( $\alpha$ -CTLA-4).

Review of immune checkpoint inhibitors in colorectal cancer

Blockade of the PD-1 (Programmed Death-1) ligand with monoclonal antibody as shows efficacy in colorectal cancer, especially those deficient in mismatch repair protein expression, as manifested by microsatellite instability (MSI-High) on pathologic exam.

Keynote-177 trial (André et al., 2020) randomized 307 patients with MSI-High dMMR colorectal cancer patients to receive either pembrolizumab or 5-FU based chemotherapy and found superior survival with pembro.



## Chapter 38

# Colectomy

### 38.1 Extended Node dissection

Short-term outcomes of complete mesocolic excision versus D2 dissection in patients undergoing laparoscopic colectomy for right colon cancer (RELARC): a randomised, controlled, phase 3, superiority trial

Short-term outcomes of a multicentre randomized clinical trial comparing D2 versus D3 lymph node dissection for colonic cancer (COLD trial). Karachun A, Panaiotti L, Chernikovskiy I, Achkasov S, Gevorkyan Y, Savanovich N, Sharygin G, Markushin L, Sushkov O, Aleshin D, Shakhmatov D, Nazarov I, Muratov I, Maynovskaya O, Olkina A, Lankov T, Ovchinnikova T, Kharagezov D, Kaymakchi D, Milakin A, Petrov A. *Br J Surg.* 2020 Apr;107(5):499-508. doi: 10.1002/bjs.11387. Epub 2019 Dec 24. PMID: 31872869 Clinical Trial.



## Chapter 39

# Chemotherapy

### 39.1 Neoadjuvant Chemotherapy

Seymour MT, Morton D. FOxTROT: an international randomized controlled trial in 1052 patients (pts) evaluating neoadjuvant chemotherapy (NAC) for colon cancer. J Clin Oncol. 2019 May;37(15 Suppl):3504-3504.





# Rectal Cancer



## Chapter 40

# Rectal Overview

### Resources

ASCRS Guidelines for treatment of Colorectal Cancer 2022



## Chapter 41

# Rectal Cancer Staging

### 41.1 MRI staging

MRI has supplanted endoscopic ultrasound (EUS) as the study of choice for staging rectal cancer.

MERCURY group trial demonstrated the predictive value of MRI for rectal cancer (Taylor et al., 2014) (MERCURY Study Group, 2006)



## Chapter 42

# Rectal Cancer Surgery

Importance of total mesorectal excision was championed by Bill Heald (Heald and Ryall, 1986) who emphasized sharp dissection of the mesorecum outside of the visceral fascial envelope. In addition he advocated dissection of the totality of the mesorectum distal to the tumor to avoid leaving behind nodes within the mesorectum distal to the tumor (Quirke et al., 1986) (Nagtegaal and Quirke, 2008) (Paty et al., 1994).





## Chapter 43

# Rectal Adjuvant Therapy

German Rectal Cancer Study (AIO-94) compared preop and postop chemoradiation in locally-advanced rectal cancer (Sauer et al., 2001). Among 823 patients, local recurrence was 6% in the preoperative group vs 13% in the postop group ( $p=0.0006$ ). Overall survival was not different, but preop chemoradiation was less toxic. Pathologic complete response rate in the preoperative group as 8%.

NSABP R-03 clinical trial (Roh et al., 2009) randomized 267 patients with T3 or T4 or node-positive rectal cancer to preop vs postop chemoradiation. Disease-free survival was better in the preoperative group, wiht no difference in overall survival. Pathologic complete response rate in the preop group was 15%.

Dutch Rectal Cancer Trial (CKVO 95-04): Demonstrated the benefit of preop radiation in combination with TME(Kapiteijn et al., 2001). Radiation reduced local recurrence form 10.9% to 5.6%

### 43.0.1 Short-course preoperative therapy

Swedish Rectal Cancer Trial compared surgery alone with preoperative short-course therapy consisting of 500cGy of radiation without chemotherapy administered in one week prior to surgery. Local recurrence was 9% in the therapy group vs 26% in the control group, with an improvement in overall survival of 38% vs 80%(Swedish Rectal Cancer Trial et al., 1997). Of note, this trial was performed in the era prioro to the widespread use of total mesorectal excision.

Stockholm III trial showed that short-course radiation therapy performed 4-8 weeks prior to surgery resulted in imporved rates of pathologic complete response (12% vs 2%) compared with short-course radiation therapy performed the week prior to surgery (Pettersson et al., 2015)

The Dutch trial found a reduction in local recurrence form 10.9% to 5.6% with preoperative radiation with no difference in survival(Kapiteijn et al., 2001). A

later update (van Gijn et al., 2011) The Medical Research Council C07 trial compared preoperative short-course radiation with selective post-operative chemoradiation and found no difference in local recurrence (4.4% vs 10.6%) and no difference in overall survival. (Sebag-Montefiore et al., 2009)

### 43.0.2 Total Neoadjuvant therapy

RAPIDO trial randomized 920 patients with T4 or node-positive disease to long-course chemoradiation followed by surgery vs short-course radiation followed by chemotherapy and surgery. The pCR rate was significantly higher in the short course/chemotherapy/surgery group (28% vs 14%) and disease-specific survival at 3 years was higher (30% vs 24%). (Bahadoer et al., 2021) (van der Valk et al., 2020)

PRODIGE 23 randomized 461 patients with T3 or T4 recta cancers to long-course radiation followed by surgery vs induction chemotherapy, long-course radiation followed by surgery. Up-front chemotherapy was associated with increased 3-year survival (76% vs 69%) and an increase in rate of pathologic complete response of 28% vs 12%. (Conroy et al., 2021)

STELLAR trial (Jin et al., 2022) TNT with short-course radiation (5Gy x 5 days) + CAPEOX chemotherapy vs long-course chemoRT with capecitabine. No differences in relapse-free survival or metastasis free survival. Overall survival better with TNT ( $p=0.03$ ). On subgroup analysis, no group appeared to have more benefit. pCR rate 17% with TNT vs 13.9% with conventional chemoradiation. Positive margins (R1) occurred in 8.5% of TNT patients vs 12.5% in the conventional chemoRT group.

OPRA clinical trial (Garcia-Aguilar et al., 2022). 324 rectal cancer patients staged with MRI randomized to INCT (induction chemo followed by chemoRT) vs CNCT (Chemoradiation followed by chemotherapy). Patients with a response were offered watch and wait. Patients without a response were treated with surgery. No difference in disease-free survival or overall survival. Among patients in watch and wait, somewhat more patients with INCT had recurrences (42 patients vs 32 patients). More organ preservation with CNCT. See also (Smith et al., 2015)

## 43.1 Neoadjuvant immunotherapy (MSI high)

MSKCC trial of neoadjuvant PD-1 blockade with 6 months of dostarlimab for 12 patients with MMR-deficient (MSI-high) rectal cancer resulted in 100% clinical response rate (Cercek et al., 2022). No patients were subsequently treated with either chemoradiation or surgery as originally planned.

PICC trial (Hu et al., 2022) Chinese trial of PD-1 blockade with neoadjuvant toripalimab + colecoxib vs toripalimab for 3 months in 34 patients with MMR-deficient locally-advanced (T3/4 or N+) colorectal cancer. All patients were then treated with surgery. Pathologic complete response in 88% of dual-therapy patients and 65% of monotherapy. Grade 3 toxicity in 1/34 patients. Previous neoadjuvant chemo in 25%. Rectal cancer in 18%.

## **43.2 Non-operative management of Rectal Cancer**

MSKCC published an experience of 113 patients with cCR after chemoradiation for rectal cancer who elected non-operative management (Smith et al., 2019). Among this group, 22 developed a local recurrence, and 8% developed metastatic disease. (Smith et al., 2020)



# Sarcoma



## Chapter 44

# Soft Tissue Sarcomas

### 44.1 Desmoid Tumors

Review of desmoid tumors: Global Consensus Guidelines

### 44.2 Retroperitoneal

#### **Preop Radiation**

STRASS trial randomized 266 patients to preop radiation followed by surgical resection vs surgery alone. At a median followup of 43 months, there was no difference in recurrence-free survival. Serious adverse affects were more common in the preop radiation group (24% vs 10%). One patients in the radiation group died of treatment-related toxicity (gastrocolic fistula), compared with none in the surgery alone group (Bonvalot et al., 2020). See commentary: (Cardona, 2020)

STRASS2: Ongoing trial of neoadjuvant chemotherapy.

### 44.3 Peritoneal mesothelioma

Review (?)

Surgical Oncology Clinics review (?)





# Bibliography

Ajani, J. A., Winter, K., Okawara, G. S., Donohue, J. H., Pisters, P. W. T., Crane, C. H., Greskovich, J. F., Anne, P. R., Bradley, J. D., Willett, C., and Rich, T. A. (2006). Phase II trial of preoperative chemoradiation in patients with localized gastric adenocarcinoma (RTOG 9904): Quality of combined modality therapy and pathologic response. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 24(24):3953–3958.

Al-Batran, S.-E., Homann, N., Pauligk, C., Goetze, T. O., Meiler, J., Kasper, S., Kopp, H.-G., Mayer, F., Haag, G. M., Luley, K., Lindig, U., Schmiegel, W., Pohl, M., Stoecklacher, J., Folprecht, G., Probst, S., Prasnikar, N., Fischbach, W., Mahlberg, R., Trojan, J., Koenigsmann, M., Martens, U. M., Thuss-Patience, P., Egger, M., Block, A., Heinemann, V., Illerhaus, G., Moehler, M., Schenk, M., Kullmann, F., Behringer, D. M., Heike, M., Pink, D., Teschendorf, C., Löhr, C., Bernhard, H., Schuch, G., Rethwisch, V., von Weikersthal, L. F., Hartmann, J. T., Kneba, M., Daum, S., Schulmann, K., Weniger, J., Belle, S., Gaiser, T., Oduncu, F. S., Güntner, M., Hozaeel, W., Reichart, A., Jäger, E., Kraus, T., Mönig, S., Bechstein, W. O., Schuler, M., Schmalenberg, H., Hofheinz, R. D., and FLOT4-AIO Investigators (2019). Perioperative chemotherapy with fluorouracil plus leucovorin, oxaliplatin, and docetaxel versus fluorouracil or capecitabine plus cisplatin and epirubicin for locally advanced, resectable gastric or gastro-oesophageal junction adenocarcinoma (FLOT4): A randomised, phase 2/3 trial. *Lancet (London, England)*, 393(10184):1948–1957.

André, T., Shiu, K.-K., Kim, T. W., Jensen, B. V., Jensen, L. H., Punt, C., Smith, D., Garcia-Carbonero, R., Benavides, M., Gibbs, P., de la Fouchardiere, C., Rivera, F., Elez, E., Bendell, J., Le, D. T., Yoshino, T., Van Cutsem, E., Yang, P., Farooqui, M. Z. H., Marinello, P., Diaz, L. A., and KEYNOTE-177 Investigators (2020). Pembrolizumab in Microsatellite-Instability-High Advanced Colorectal Cancer. *The New England Journal of Medicine*, 383(23):2207–2218.

Bahadoer, R. R., Dijkstra, E. A., van Etten, B., Marijnen, C. A. M., Putter, H., Kranenbarg, E. M.-K., Roodvoets, A. G. H., Nagtegaal, I. D., Beets-Tan,

- R. G. H., Blomqvist, L. K., Fokstuen, T., Ten Tije, A. J., Capdevila, J., Hendriks, M. P., Edhemovic, I., Cervantes, A., Nilsson, P. J., Glimelius, B., van de Velde, C. J. H., Hospers, G. A. P., and RAPIDO collaborative investigators (2021). Short-course radiotherapy followed by chemotherapy before total mesorectal excision (TME) versus preoperative chemoradiotherapy, TME, and optional adjuvant chemotherapy in locally advanced rectal cancer (RAPIDO): A randomised, open-label, phase 3 trial. *The Lancet. Oncology*, 22(1):29–42.
- Baker, M., Halliday, V., Williams, R. N., and Bowrey, D. J. (2016). A systematic review of the nutritional consequences of esophagectomy. *Clinical Nutrition (Edinburgh, Scotland)*, 35(5):987–994.
- Bang, Y.-J., Kim, Y.-W., Yang, H.-K., Chung, H. C., Park, Y.-K., Lee, K. H., Lee, K.-W., Kim, Y. H., Noh, S.-I., Cho, J. Y., Mok, Y. J., Kim, Y. H., Ji, J., Yeh, T.-S., Button, P., Sirzén, F., Noh, S. H., and CLASSIC trial investigators (2012). Adjuvant capecitabine and oxaliplatin for gastric cancer after D2 gastrectomy (CLASSIC): A phase 3 open-label, randomised controlled trial. *Lancet (London, England)*, 379(9813):315–321.
- Bedenne, L., Michel, P., Bouché, O., Milan, C., Mariette, C., Conroy, T., Pezet, D., Roulet, B., Seitz, J.-F., Herr, J.-P., Paillot, B., Arveux, P., Bonnetain, F., and Binequet, C. (2007). Chemoradiation followed by surgery compared with chemoradiation alone in squamous cancer of the esophagus: FFCO 9102. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 25(10):1160–1168.
- Bergeron, E. J., Lin, J., Chang, A. C., Orringer, M. B., and Reddy, R. M. (2014). Endoscopic ultrasound is inadequate to determine which T1/T2 esophageal tumors are candidates for endoluminal therapies. *The Journal of Thoracic and Cardiovascular Surgery*, 147(2):765–771: Discussion 771–773.
- Beukema, J. C., van Luijk, P., Widder, J., Langendijk, J. A., and Muijs, C. T. (2015). Is cardiac toxicity a relevant issue in the radiation treatment of esophageal cancer? *Radiotherapy and Oncology: Journal of the European Society for Therapeutic Radiology and Oncology*, 114(1):85–90.
- Biere, S. S., van Berge Henegouwen, M. I., Maas, K. W., Bonavina, L., Rosman, C., Garcia, J. R., Gisbertz, S. S., Klinkenbijl, J. H., Hollmann, M. W., de Lange, E. S., Bonjer, H. J., van der Peet, D. L., and Cuesta, M. A. (2012). Minimally invasive versus open oesophagectomy for patients with oesophageal cancer: A multicentre, open-label, randomised controlled trial. *Lancet*, 379(9829):1887–92.
- Birkmeyer, J. D., Stukel, T. A., Siewers, A. E., Goodney, P. P., Wennberg, D. E., and Lucas, F. L. (2003). Surgeon volume and operative mortality in the United States. *The New England Journal of Medicine*, 349(22):2117–2127.

- Bonvalot, S., Gronchi, A., Le Péchoux, C., Swallow, C. J., Strauss, D., Meeus, P., van Coevorden, F., Stoldt, S., Stoeckle, E., Rutkowski, P., Rastrelli, M., Raut, C. P., Hompes, D., De Paoli, A., Sangalli, C., Honoré, C., Chung, P., Miah, A., Blay, J. Y., Fiore, M., Stelmes, J.-J., Dei Tos, A. P., Baldini, E. H., Litière, S., Marreaud, S., Gelderblom, H., and Haas, R. L. (2020). Preoperative radiotherapy plus surgery versus surgery alone for patients with primary retroperitoneal sarcoma (EORTC-62092: STRASS): A multicentre, open-label, randomised, phase 3 trial. *The Lancet. Oncology*, 21(10):1366–1377.
- Cao, Y., Liao, C., Tan, A., Gao, Y., Mo, Z., and Gao, F. (2009). Meta-analysis of endoscopic submucosal dissection versus endoscopic mucosal resection for tumors of the gastrointestinal tract. *Endoscopy*, 41(9):751–757.
- Cardona, K. (2020). The STRASS trial: An important step in the right direction. *The Lancet. Oncology*, 21(10):1257–1258.
- Castoro, C., Scarpa, M., Cagol, M., Alfieri, R., Ruol, A., Cavallin, F., Michieletto, S., Zanchettin, G., Chiarion-Sileni, V., Corti, L., and Ancona, E. (2013). Complete clinical response after neoadjuvant chemoradiotherapy for squamous cell cancer of the thoracic oesophagus: Is surgery always necessary? *Journal of Gastrointestinal Surgery: Official Journal of the Society for Surgery of the Alimentary Tract*, 17(8):1375–1381.
- Cercek, A., Lumish, M., Sinopoli, J., Weiss, J., Shia, J., Lamendola-Essel, M., El Dika, I. H., Segal, N., Shcherba, M., Sugarman, R., Stadler, Z., Yaeger, R., Smith, J. J., Rousseau, B., Argiles, G., Patel, M., Desai, A., Saltz, L. B., Widmar, M., Iyer, K., Zhang, J., Gianino, N., Crane, C., Romesser, P. B., Pappou, E. P., Paty, P., Garcia-Aguilar, J., Gonen, M., Gollub, M., Weiser, M. R., Schalper, K. A., and Diaz, L. A. (2022). PD-1 Blockade in Mismatch Repair-Deficient, Locally Advanced Rectal Cancer. *The New England Journal of Medicine*, 386(25):2363–2376.
- Chatterton, B. E., Ho Shon, I., Baldey, A., Lenzo, N., Patrikeos, A., Kelley, B., Wong, D., Ramshaw, J. E., and Scott, A. M. (2009). Positron emission tomography changes management and prognostic stratification in patients with oesophageal cancer: Results of a multicentre prospective study. *European Journal of Nuclear Medicine and Molecular Imaging*, 36(3):354–361.
- Chirieac, L. R., Swisher, S. G., Ajani, J. A., Komaki, R. R., Correa, A. M., Morris, J. S., Roth, J. A., Rashid, A., Hamilton, S. R., and Wu, T.-T. (2005). Posttherapy pathologic stage predicts survival in patients with esophageal carcinoma receiving preoperative chemoradiation. *Cancer*, 103(7):1347–1355.
- Conroy, T., Bosset, J.-F., Etienne, P.-L., Rio, E., François, É., Mesgouez-Nebout, N., Vendrely, V., Artignan, X., Bouché, O., Gargot, D., Boige, V., Bonichon-Lamichhane, N., Louvet, C., Morand, C., de la Fouchardière, C., Lamfichekh, N., Juzyna, B., Jouffroy-Zeller, C., Rullier, E., Marchal, F.,

- Gourgou, S., Castan, F., Borg, C., and Unicancer Gastrointestinal Group and Partenariat de Recherche en Oncologie Digestive (PRODIGE) Group (2021). Neoadjuvant chemotherapy with FOLFIRINOX and preoperative chemoradiotherapy for patients with locally advanced rectal cancer (UNICANCER-PRODIGE 23): A multicentre, randomised, open-label, phase 3 trial. *The Lancet. Oncology*, 22(5):702–715.
- Cunningham, D., Allum, W. H., Stenning, S. P., Thompson, J. N., Van de Velde, C. J. H., Nicolson, M., Scarffe, J. H., Lofts, F. J., Falk, S. J., Iveson, T. J., Smith, D. B., Langley, R. E., Verma, M., Weeden, S., Chua, Y. J., and MAGIC Trial Participants, n. (2006). Perioperative chemotherapy versus surgery alone for resectable gastroesophageal cancer. *The New England Journal of Medicine*, 355(1):11–20.
- de Graaf, G. W., Ayantunde, A. A., Parsons, S. L., Duffy, J. P., and Welch, N. T. (2007). The role of staging laparoscopy in oesophagogastric cancers. *European Journal of Surgical Oncology: The Journal of the European Society of Surgical Oncology and the British Association of Surgical Oncology*, 33(8):988–992.
- Dorth, J. A., Pura, J. A., Palta, M., Willett, C. G., Uronis, H. E., D’Amico, T. A., and Czito, B. G. (2014). Patterns of recurrence after trimodality therapy for esophageal cancer. *Cancer*, 120(14):2099–2105.
- Dunbar, K. B. and Spechler, S. J. (2012). The risk of lymph-node metastases in patients with high-grade dysplasia or intramucosal carcinoma in Barrett’s esophagus: A systematic review. *The American Journal of Gastroenterology*, 107(6):850–862; quiz 863.
- Frandsen, J., Boothe, D., Gaffney, D. K., Wilson, B. D., and Lloyd, S. (2015). Increased risk of death due to heart disease after radiotherapy for esophageal cancer. *Journal of Gastrointestinal Oncology*, 6(5):516–523.
- Furlong, H., Bass, G., Breathnach, O., O’Neill, B., Leen, E., and Walsh, T. N. (2013). Targeting therapy for esophageal cancer in patients aged 70 and over. *Journal of Geriatric Oncology*, 4(2):107–113.
- Ganz, R. A., Overholt, B. F., Sharma, V. K., Fleischer, D. E., Shaheen, N. J., Lightdale, C. J., Freeman, S. R., Pruitt, R. E., Urayama, S. M., Gress, F., Pavey, D. A., Branch, M. S., Savides, T. J., Chang, K. J., Muthusamy, V. R., Bohorfoush, A. G., Pace, S. C., DeMeester, S. R., Eysselein, V. E., Panjehpour, M., Triadafilopoulos, G., and U.S. Multicenter Registry (2008). Circumferential ablation of Barrett’s esophagus that contains high-grade dysplasia: A U.S. Multicenter Registry. *Gastrointestinal Endoscopy*, 68(1):35–40.
- Garcia-Aguilar, J., Patil, S., Gollub, M. J., Kim, J. K., Yuval, J. B., Thompson, H. M., Verheij, F. S., Omer, D. M., Lee, M., Dunne, R. F., Marcet, J., Cataldo, P., Polite, B., Herzig, D. O., Liska, D., Oommen, S., Friel, C. M., Ternent, C., Coveler, A. L., Hunt, S., Gregory, A., Varma, M. G., Bello, B. L., Carmichael, J. C., Krauss, J., Gleisner, A., Paty, P. B., Weiser, M. R., Nash,

- G. M., Pappou, E., Guillem, J. G., Temple, L., Wei, I. H., Widmar, M., Lin, S., Segal, N. H., Cercek, A., Yaeger, R., Smith, J. J., Goodman, K. A., Wu, A. J., and Saltz, L. B. (2022). Organ Preservation in Patients With Rectal Adenocarcinoma Treated With Total Neoadjuvant Therapy. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, page JCO2200032.
- Gharzai, L., Verma, V., Denniston, K. A., Bhirud, A. R., Bennion, N. R., and Lin, C. (2016). Radiation Therapy and Cardiac Death in Long-Term Survivors of Esophageal Cancer: An Analysis of the Surveillance, Epidemiology, and End Result Database. *PLoS One*, 11(7):e0158916.
- Guttmann, D. M., Mitra, N., Bekelman, J., Metz, J. M., Plataras, J., Feng, W., and Swisher-McClure, S. (2017). Improved Overall Survival with Aggressive Primary Tumor Radiotherapy for Patients with Metastatic Esophageal Cancer. *Journal of Thoracic Oncology: Official Publication of the International Association for the Study of Lung Cancer*, 12(7):1131–1142.
- Haidry, R. J., Dunn, J. M., Butt, M. A., Burnell, M. G., Gupta, A., Green, S., Miah, H., Smart, H. L., Bhandari, P., Smith, L. A., Willert, R., Fullarton, G., Morris, J., Di Pietro, M., Gordon, C., Penman, I., Barr, H., Patel, P., Boger, P., Kapoor, N., Mahon, B., Hoare, J., Narayanasamy, R., O’Toole, D., Cheong, E., Direkze, N. C., Ang, Y., Novelli, M., Banks, M. R., and Lovat, L. B. (2013). Radiofrequency ablation and endoscopic mucosal resection for dysplastic barrett’s esophagus and early esophageal adenocarcinoma: Outcomes of the UK National Halo RFA Registry. *Gastroenterology*, 145(1):87–95.
- Heald, R. J. and Ryall, R. D. (1986). Recurrence and survival after total mesorectal excision for rectal cancer. *Lancet (London, England)*, 1(8496):1479–1482.
- Hollis, A. C., Quinn, L. M., Hodson, J., Evans, E., Plowright, J., Begum, R., Mitchell, H., Hallissey, M. T., Whiting, J. L., and Griffiths, E. A. (2017). Prognostic significance of tumor length in patients receiving esophagectomy for esophageal cancer. *Journal of Surgical Oncology*, 116(8):1114–1122.
- Hu, H., Kang, L., Zhang, J., Wu, Z., Wang, H., Huang, M., Lan, P., Wu, X., Wang, C., Cao, W., Hu, J., Huang, Y., Huang, L., Wang, H., Shi, L., Cai, Y., Shen, C., Ling, J., Xie, X., Cai, Y., He, X., Dou, R., Zhou, J., Ma, T., Zhang, X., Luo, S., Deng, W., Ling, L., Liu, H., and Deng, Y. (2022). Neoadjuvant PD-1 blockade with toripalimab, with or without celecoxib, in mismatch repair-deficient or microsatellite instability-high, locally advanced, colorectal cancer (PICC): A single-centre, parallel-group, non-comparative, randomised, phase 2 trial. *The Lancet. Gastroenterology & Hepatology*, 7(1):38–48.
- Hulscher, J. B. F., van Sandick, J. W., de Boer, A. G. E. M., Wijnhoven, B. P. L., Tijssen, J. G. P., Fockens, P., Stalmeier, P. F. M., ten Kate, F. J. W.,

- van Dekken, H., Obertop, H., Tilanus, H. W., and van Lanschot, J. J. B. (2002). Extended transthoracic resection compared with limited transhiatal resection for adenocarcinoma of the esophagus. *The New England Journal of Medicine*, 347(21):1662–1669.
- Jin, J., Tang, Y., Hu, C., Jiang, L.-M., Jiang, J., Li, N., Liu, W.-Y., Chen, S.-L., Li, S., Lu, N.-N., Cai, Y., Li, Y.-H., Zhu, Y., Cheng, G.-H., Zhang, H.-Y., Wang, X., Zhu, S.-Y., Wang, J., Li, G.-F., Yang, J.-L., Zhang, K., Chi, Y., Yang, L., Zhou, H.-T., Zhou, A.-P., Zou, S.-M., Fang, H., Wang, S.-L., Zhang, H.-Z., Wang, X.-S., Wei, L.-C., Wang, W.-L., Liu, S.-X., Gao, Y.-H., and Li, Y.-X. (2022). Multicenter, Randomized, Phase III Trial of Short-Term Radiotherapy Plus Chemotherapy Versus Long-Term Chemoradiotherapy in Locally Advanced Rectal Cancer (STELLAR). *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 40(15):1681–1692.
- Kalff, M. C., Fransen, L. F. C., de Groot, E. M., Gisbertz, S. S., Nieuwenhuijzen, G. A. P., Ruurda, J. P., Verhoeven, R. H. A., Luyer, M. D. P., van Hillegersberg, R., van Berge Henegouwen, M. I., and Dutch Upper Gastrointestinal Cancer Audit group (2020). Long-Term Survival After Minimally Invasive Versus Open Esophagectomy for Esophageal Cancer: A Nationwide Propensity-Score Matched Analysis. *Annals of Surgery*, Publish Ahead of Print.
- Kapiteijn, E., Marijnen, C. A., Nagtegaal, I. D., Putter, H., Steup, W. H., Wiggers, T., Rutten, H. J., Pahlman, L., Glimelius, B., van Krieken, J. H., Leer, J. W., van de Velde, C. J., and Dutch Colorectal Cancer Group (2001). Pre-operative radiotherapy combined with total mesorectal excision for resectable rectal cancer. *The New England Journal of Medicine*, 345(9):638–646.
- Kato, H., Kuwano, H., Nakajima, M., Miyazaki, T., Yoshikawa, M., Ojima, H., Tsukada, K., Oriuchi, N., Inoue, T., and Endo, K. (2002). Comparison between positron emission tomography and computed tomography in the use of the assessment of esophageal carcinoma. *Cancer*, 94(4):921–928.
- Khan, Q. J. and Fabian, C. J. (2010). How I Treat Vitamin D Deficiency. *Journal of Oncology Practice*, 6(2):97–101.
- Kim, T. J., Kim, H. Y., Lee, K. W., and Kim, M. S. (2009 Mar-Apr). Multimodality assessment of esophageal cancer: Preoperative staging and monitoring of response to therapy. *Radiographics: A Review Publication of the Radiological Society of North America, Inc*, 29(2):403–421.
- Kofoed, S. C., Muhic, A., Baeksgaard, L., Jendresen, M., Gustafsen, J., Holm, J., Bardram, L., Brandt, B., Brenø, J., and Svendsen, L. B. (2012). Survival after adjuvant chemoradiotherapy or surgery alone in resectable adenocarcinoma at the gastro-esophageal junction. *Scandinavian journal of surgery: SJS: official organ for the Finnish Surgical Society and the Scandinavian Surgical Society*, 101(1):26–31.

- Lee, J., Lim, D. H., Kim, S., Park, S. H., Park, J. O., Park, Y. S., Lim, H. Y., Choi, M. G., Sohn, T. S., Noh, J. H., Bae, J. M., Ahn, Y. C., Sohn, I., Jung, S. H., Park, C. K., Kim, K.-M., and Kang, W. K. (2012). Phase III trial comparing capecitabine plus cisplatin versus capecitabine plus cisplatin with concurrent capecitabine radiotherapy in completely resected gastric cancer with D2 lymph node dissection: The ARTIST trial. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 30(3):268–273.
- Leichman, L. P., Goldman, B. H., Bohanes, P. O., Lenz, H. J., Thomas, C. R., Billingsley, K. G., Corless, C. L., Iqbal, S., Gold, P. J., Benedetti, J. K., Danenberg, K. D., and Blanke, C. D. (2011). S0356: A phase II clinical and prospective molecular trial with oxaliplatin, fluorouracil, and external-beam radiation therapy before surgery for patients with esophageal adenocarcinoma. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 29(34):4555–4560.
- Macdonald, J. S., Smalley, S. R., Benedetti, J., Hundahl, S. A., Estes, N. C., Stemmermann, G. N., Haller, D. G., Ajani, J. A., Gunderson, L. L., Jessup, J. M., and Martenson, J. A. (2001). Chemoradiotherapy after surgery compared with surgery alone for adenocarcinoma of the stomach or gastroesophageal junction. *The New England Journal of Medicine*, 345(10):725–730.
- Maish, M. S. and DeMeester, S. R. (2004). Endoscopic mucosal resection as a staging technique to determine the depth of invasion of esophageal adenocarcinoma. *The Annals of Thoracic Surgery*, 78(5):1777–1782.
- Mariette, C., Markar, S. R., Dabakuyo-Yonli, T. S., Meunier, B., Pezet, D., Collet, D., D'Journo, X. B., Brigand, C., Perniceni, T., Carrère, N., Mabrut, J.-Y., Msika, S., Peschaud, F., Prudhomme, M., Bonnetain, F., and Piessen, G. (2019). Hybrid Minimally Invasive Esophagectomy for Esophageal Cancer. *New England Journal of Medicine*, 380(2):152–162.
- Markar, S. R., Gronnier, C., Pasquer, A., Duhamel, A., Beal, H., Théreaux, J., Gagnière, J., Lebreton, G., Brigand, C., Meunier, B., Collet, D., Mariette, C., and FREGAT working group – FRENCH – AFC (2016). Role of neoadjuvant treatment in clinical T2N0M0 oesophageal cancer: Results from a retrospective multi-center European study. *European Journal of Cancer (Oxford, England: 1990)*, 56:59–68.
- Markar, S. R., Karthikesalingam, A., Penna, M., and Low, D. E. (2014). Assessment of short-term clinical outcomes following salvage esophagectomy for the treatment of esophageal malignancy: Systematic review and pooled analysis. *Annals of Surgical Oncology*, 21(3):922–931.
- Martin, L. and Lagergren, P. (2009). Long-term weight change after oesophageal cancer surgery. *The British Journal of Surgery*, 96(11):1308–1314.
- McKeown, K. C. (1976). Total three-stage oesophagectomy for cancer of the oesophagus. *The British Journal of Surgery*, 63(4):259–262.

- Mellow, M. H. and Pinkas, H. (1985). Endoscopic laser therapy for malignancies affecting the esophagus and gastroesophageal junction. Analysis of technical and functional efficacy. *Archives of Internal Medicine*, 145(8):1443–1446.
- MERCURY Study Group (2006). Diagnostic accuracy of preoperative magnetic resonance imaging in predicting curative resection of rectal cancer: Prospective observational study. *BMJ (Clinical research ed.)*, 333(7572):779.
- Messenger, M., Mirabel, X., Tresch, E., Paumier, A., Vendrely, V., Dahan, L., Glehen, O., Vasseur, F., Lacornerie, T., Piessen, G., El Hajbi, F., Robb, W. B., Clisant, S., Kramar, A., Mariette, C., and Adenis, A. (2016). Pre-operative chemoradiation with paclitaxel-carboplatin or with fluorouracil-oxaliplatin-folinic acid (FOLFOX) for resectable esophageal and junctional cancer: The PROTECT-1402, randomized phase 2 trial. *BMC cancer*, 16:318.
- Minsky, B. D., Pajak, T. F., Ginsberg, R. J., Pisansky, T. M., Martenson, J., Komaki, R., Okawara, G., Rosenthal, S. A., and Kelsen, D. P. (2002). INT 0123 (Radiation Therapy Oncology Group 94-05) phase III trial of combined-modality therapy for esophageal cancer: High-dose versus standard-dose radiation therapy. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 20(5):1167–1174.
- Nagtegaal, I. D. and Quirke, P. (2008). What is the role for the circumferential margin in the modern treatment of rectal cancer? *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 26(2):303–312.
- Noh, S. H., Park, S. R., Yang, H.-K., Chung, H. C., Chung, I.-J., Kim, S.-W., Kim, H.-H., Choi, J.-H., Kim, H.-K., Yu, W., Lee, J. I., Shin, D. B., Ji, J., Chen, J.-S., Lim, Y., Ha, S., Bang, Y.-J., and CLASSIC trial investigators (2014). Adjuvant capecitabine plus oxaliplatin for gastric cancer after D2 gastrectomy (CLASSIC): 5-year follow-up of an open-label, randomised phase 3 trial. *The Lancet. Oncology*, 15(12):1389–1396.
- Nurkin, S. J., Nava, H. R., Yendamuri, S., LeVea, C. M., Nwogu, C. E., Groman, A., Wilding, G., Bain, A. J., Hochwald, S. N., and Khushalani, N. I. (2014). Outcomes of endoscopic resection for high-grade dysplasia and esophageal cancer. *Surgical Endoscopy*, 28(4):1090–1095.
- Omloo, J. M. T., Lagarde, S. M., Hulscher, J. B. F., Reitsma, J. B., Fockens, P., van Dekken, H., Ten Kate, F. J. W., Obertop, H., Tilanus, H. W., and van Lanschot, J. J. B. (2007). Extended transthoracic resection compared with limited transhiatal resection for adenocarcinoma of the mid/distal esophagus: Five-year survival of a randomized clinical trial. *Annals of Surgery*, 246(6):992–1000; discussion 1000–1001.
- Oppedijk, V., van der Gaast, A., van Lanschot, J. J. B., van Hagen, P., van Os, R., van Rij, C. M., van der Sangen, M. J., Beukema, J. C., Rütten, H., Spruit, P. H., Reinders, J. G., Richel, D. J., van Berge Henegouwen, M. I., and



- Hulshof, M. C. C. M. (2014). Patterns of recurrence after surgery alone versus preoperative chemoradiotherapy and surgery in the CROSS trials. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 32(5):385–391.
- Orringer, M. B., Marshall, B., Chang, A. C., Lee, J., Pickens, A., and Lau, C. L. (2007). Two thousand transhiatal esophagectomies: Changing trends, lessons learned. *Annals of Surgery*, 246(3):363–372; discussion 372–374.
- Orringer, M. B. and Sloan, H. (1978). Esophagectomy without thoracotomy. *The Journal of Thoracic and Cardiovascular Surgery*, 76(5):643–654.
- Ouattara, M., D’Journo, X. B., Loundou, A., Trousse, D., Dahan, L., Doddoli, C., Seitz, J. F., and Thomas, P.-A. (2012). Body mass index kinetics and risk factors of malnutrition one year after radical oesophagectomy for cancer. *European Journal of Cardio-Thoracic Surgery: Official Journal of the European Association for Cardio-Thoracic Surgery*, 41(5):1088–1093.
- Pasquali, S., Yim, G., Vohra, R. S., Mocellin, S., Nyanhongo, D., Marriott, P., Geh, J. I., and Griffiths, E. A. (2017). Survival After Neoadjuvant and Adjuvant Treatments Compared to Surgery Alone for Resectable Esophageal Carcinoma: A Network Meta-analysis. *Annals of Surgery*, 265(3):481–491.
- Paty, P. B., Enker, W. E., Cohen, A. M., and Lauwers, G. Y. (1994). Treatment of rectal cancer by low anterior resection with coloanal anastomosis. *Annals of Surgery*, 219(4):365–373.
- Pech, O., Behrens, A., May, A., Nachbar, L., Gossner, L., Rabenstein, T., Manner, H., Guenter, E., Huijsmans, J., Vieth, M., Stolte, M., and Ell, C. (2008). Long-term results and risk factor analysis for recurrence after curative endoscopic therapy in 349 patients with high-grade intraepithelial neoplasia and mucosal adenocarcinoma in Barrett’s oesophagus. *Gut*, 57(9):1200–1206.
- Pech, O., May, A., Manner, H., Behrens, A., Pohl, J., Weferling, M., Hartmann, U., Manner, N., Huijsmans, J., Gossner, L., Rabenstein, T., Vieth, M., Stolte, M., and Ell, C. (2014). Long-term efficacy and safety of endoscopic resection for patients with mucosal adenocarcinoma of the esophagus. *Gastroenterology*, 146(3):652–660.e1.
- Penniment, M. G., De Ieso, P. B., Harvey, J. A., Stephens, S., Au, H.-J., O’Callaghan, C. J., Kneebone, A., Ngan, S. Y., Ward, I. G., Roy, R., Smith, J. G., Nijjar, T., Biagi, J. J., Mulroy, L. A., Wong, R., and TROG 03.01/CCTG ES.2 group (2018). Palliative chemoradiotherapy versus radiotherapy alone for dysphagia in advanced oesophageal cancer: A multicentre randomised controlled trial (TROG 03.01). *The Lancet. Gastroenterology & Hepatology*, 3(2):114–124.
- Pettersson, D., Löhrinc, E., Holm, T., Iversen, H., Cedermarck, B., Glimelius, B., and Martling, A. (2015). Tumour regression in the randomized Stockholm

- III Trial of radiotherapy regimens for rectal cancer. *The British Journal of Surgery*, 102(8):972–978; discussion 978.
- Phoa, K. N., van Vilsteren, F. G. I., Weusten, B. L. A. M., Bisschops, R., Schoon, E. J., Ragunath, K., Fullarton, G., Di Pietro, M., Ravi, N., Visser, M., Offerhaus, G. J., Seldenrijk, C. A., Meijer, S. L., ten Kate, F. J. W., Tijssen, J. G. P., and Bergman, J. J. G. H. M. (2014). Radiofrequency ablation vs endoscopic surveillance for patients with Barrett esophagus and low-grade dysplasia: A randomized clinical trial. *JAMA*, 311(12):1209–1217.
- Poultides, G. A. and Paty, P. B. (2011). Reassessing the need for primary tumor surgery in unresectable metastatic colorectal cancer: Overview and perspective. *Therapeutic Advances in Medical Oncology*, 3(1):35–42.
- Quirke, P., Durdey, P., Dixon, M. F., and Williams, N. S. (1986). Local recurrence of rectal adenocarcinoma due to inadequate surgical resection. Histopathological study of lateral tumour spread and surgical excision. *Lancet (London, England)*, 2(8514):996–999.
- Repici, A., Hassan, C., Carlino, A., Pagano, N., Zullo, A., Rando, G., Strangio, G., Romeo, F., Nicita, R., Rosati, R., and Malesci, A. (2010). Endoscopic submucosal dissection in patients with early esophageal squamous cell carcinoma: Results from a prospective Western series. *Gastrointestinal Endoscopy*, 71(4):715–721.
- Rice, T. W., Mason, D. P., Murthy, S. C., Zuccaro, G., J., Adelstein, D. J., Rybicki, L. A., and Blackstone, E. H. (2007). T2N0M0 esophageal cancer. *J Thorac Cardiovasc Surg*, 133(2):317–24.
- Roh, M. S., Colangelo, L. H., O’Connell, M. J., Yothers, G., Deutsch, M., Allegra, C. J., Kahlenberg, M. S., Baez-Diaz, L., Ursiny, C. S., Petrelli, N. J., and Wolmark, N. (2009). Preoperative multimodality therapy improves disease-free survival in patients with carcinoma of the rectum: NSABP R-03. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 27(31):5124–5130.
- Rusch, V. W. (2004). Are cancers of the esophagus, gastroesophageal junction, and cardia one disease, two, or several? *Seminars in Oncology*, 31(4):444–449.
- Sarkaria, I. S., Rizk, N. P., Bains, M. S., Tang, L. H., Ilson, D. H., Minsky, B. I., and Rusch, V. W. (2009). Post-treatment endoscopic biopsy is a poor-predictor of pathologic response in patients undergoing chemoradiation therapy for esophageal cancer. *Annals of Surgery*, 249(5):764–767.
- Sauer, R., Fietkau, R., Wittekind, C., Martus, P., Rödel, C., Hohenberger, W., Jatzko, G., Sabitzer, H., Karstens, J. H., Becker, H., Hess, C., and Raab, R. (2001). Adjuvant versus neoadjuvant radiochemotherapy for locally advanced rectal cancer. A progress report of a phase-III randomized trial (protocol CAO/ARO/AIO-94). *Strahlentherapie Und Onkologie: Organ Der Deutschen Röntgengesellschaft ... [et Al]*, 177(4):173–181.

- Sebag-Montefiore, D., Stephens, R. J., Steele, R., Monson, J., Grieve, R., Khanna, S., Quirke, P., Couture, J., de Metz, C., Myint, A. S., Bessell, E., Griffiths, G., Thompson, L. C., and Parmar, M. (2009). Preoperative radiotherapy versus selective postoperative chemoradiotherapy in patients with rectal cancer (MRC CR07 and NCIC-CTG C016): A multicentre, randomised trial. *The Lancet*, 373(9666):811–820.
- Shaheen, N. J., Falk, G. W., Iyer, P. G., Gerson, L. B., and American College of Gastroenterology (2016). ACG Clinical Guideline: Diagnosis and Management of Barrett’s Esophagus. *The American Journal of Gastroenterology*, 111(1):30–50; quiz 51.
- Shaheen, N. J., Sharma, P., Overholt, B. F., Wolfsen, H. C., Sampliner, R. E., Wang, K. K., Galanko, J. A., Bronner, M. P., Goldblum, J. R., Bennett, A. E., Jobe, B. A., Eisen, G. M., Fennerty, M. B., Hunter, J. G., Fleischer, D. E., Sharma, V. K., Hawes, R. H., Hoffman, B. J., Rothstein, R. I., Gordon, S. R., Mashimo, H., Chang, K. J., Muthusamy, V. R., Edmundowicz, S. A., Spechler, S. J., Siddiqui, A. A., Souza, R. F., Infantolino, A., Falk, G. W., Kimmey, M. B., Madanick, R. D., Chak, A., and Lightdale, C. J. (2009). Radiofrequency ablation in Barrett’s esophagus with dysplasia. *The New England Journal of Medicine*, 360(22):2277–2288.
- Sharma, P., Kozarek, R., and Practice Parameters Committee of American College of Gastroenterology (2010). Role of esophageal stents in benign and malignant diseases. *The American Journal of Gastroenterology*, 105(2):258–273; quiz 274.
- Siewert, J. R., Stein, H. J., and Feith, M. (2006). Adenocarcinoma of the esophago-gastric junction. *Scandinavian journal of surgery: SJS: official organ for the Finnish Surgical Society and the Scandinavian Surgical Society*, 95(4):260–269.
- Sjoquist, K. M., Burmeister, B. H., Smithers, B. M., Zalcberg, J. R., Simes, R. J., Barbour, A., Gebiski, V., and Australasian Gastro-Intestinal Trials Group (2011). Survival after neoadjuvant chemotherapy or chemoradiotherapy for resectable oesophageal carcinoma: An updated meta-analysis. *The Lancet. Oncology*, 12(7):681–692.
- Smalley, S. R., Benedetti, J. K., Haller, D. G., Hundahl, S. A., Estes, N. C., Ajani, J. A., Gunderson, L. L., Goldman, B., Martenson, J. A., Jessup, J. M., Stemmermann, G. N., Blanke, C. D., and Macdonald, J. S. (2012). Updated analysis of SWOG-directed intergroup study 0116: A phase III trial of adjuvant radiochemotherapy versus observation after curative gastric cancer resection. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 30(19):2327–2333.
- Smith, J. J., Chow, O. S., Gollub, M. J., Nash, G. M., Temple, L. K., Weiser, M. R., Guillem, J. G., Paty, P. B., Avila, K., Garcia-Aguilar, J., and Rectal

- Cancer Consortium (2015). Organ Preservation in Rectal Adenocarcinoma: A phase II randomized controlled trial evaluating 3-year disease-free survival in patients with locally advanced rectal cancer treated with chemoradiation plus induction or consolidation chemotherapy, and total mesorectal excision or nonoperative management. *BMC cancer*, 15:767.
- Smith, J. J., Paty, P. B., and Garcia-Aguilar, J. (2020). Watch and Wait in Rectal Cancer or More Wait and See? *JAMA surgery*, 155(7):657–658.
- Smith, J. J., Strombom, P., Chow, O. S., Roxburgh, C. S., Lynn, P., Eaton, A., Widmar, M., Ganesh, K., Yaeger, R., Cercek, A., Weiser, M. R., Nash, G. M., Guillem, J. G., Temple, L. K. F., Chalasani, S. B., Fuqua, J. L., Petkovska, I., Wu, A. J., Reyngold, M., Vakiani, E., Shia, J., Segal, N. H., Smith, J. D., Crane, C., Gollub, M. J., Gonen, M., Saltz, L. B., Garcia-Aguilar, J., and Paty, P. B. (2019). Assessment of a Watch-and-Wait Strategy for Rectal Cancer in Patients With a Complete Response After Neoadjuvant Therapy. *JAMA oncology*, 5(4):e185896.
- Soetikno, R., Kaltenbach, T., Yeh, R., and Gotoda, T. (2005). Endoscopic mucosal resection for early cancers of the upper gastrointestinal tract. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 23(20):4490–4498.
- Speicher, P. J., Ganapathi, A. M., Englum, B. R., Hartwig, M. G., Onaitis, M. W., D’Amico, T. A., and Berry, M. F. (2014). Induction therapy does not improve survival for clinical stage T2N0 esophageal cancer. *Journal of Thoracic Oncology: Official Publication of the International Association for the Study of Lung Cancer*, 9(8):1195–1201.
- Stahl, M., Stuschke, M., Lehmann, N., Meyer, H.-J., Walz, M. K., Seeber, S., Klump, B., Budach, W., Teichmann, R., Schmitt, M., Schmitt, G., Franke, C., and Wilke, H. (2005). Chemoradiation with and without surgery in patients with locally advanced squamous cell carcinoma of the esophagus. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 23(10):2310–2317.
- Stahl, M., Walz, M. K., Stuschke, M., Lehmann, N., Meyer, H.-J., Riera-Knorrenschild, J., Langer, P., Engenhart-Cabillic, R., Bitzer, M., Königsrainer, A., Budach, W., and Wilke, H. (2009). Phase III comparison of preoperative chemotherapy compared with chemoradiotherapy in patients with locally advanced adenocarcinoma of the esophagogastric junction. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 27(6):851–856.
- Sudo, K., Taketa, T., Correa, A. M., Campagna, M.-C., Wadhwa, R., Blum, M. A., Komaki, R., Lee, J. H., Bhutani, M. S., Weston, B., Skinner, H. D., Maru, D. M., Rice, D. C., Swisher, S. G., Hofstetter, W. L., and Ajani, J. A. (2013). Locoregional failure rate after preoperative chemoradiation of

- esophageal adenocarcinoma and the outcomes of salvage strategies. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 31(34):4306–4310.
- Sudo, K., Xiao, L., Wadhwa, R., Shiozaki, H., Elimova, E., Taketa, T., Blum, M. A., Lee, J. H., Bhutani, M. S., Weston, B., Ross, W. A., Komaki, R., Rice, D. C., Swisher, S. G., Hofstetter, W. L., Maru, D. M., Skinner, H. D., and Ajani, J. A. (2014). Importance of surveillance and success of salvage strategies after definitive chemoradiation in patients with esophageal cancer. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 32(30):3400–3405.
- Swedish Rectal Cancer Trial, Cedermark, B., Dahlberg, M., Glimelius, B., Pahlman, L., Rutqvist, L. E., and Wilking, N. (1997). Improved survival with preoperative radiotherapy in resectable rectal cancer. *The New England Journal of Medicine*, 336(14):980–987.
- Swisher, S. G., Wynn, P., Putnam, J. B., Mosheim, M. B., Correa, A. M., Komaki, R. R., Ajani, J. A., Smythe, W. R., Vaporciyan, A. A., Roth, J. A., and Walsh, G. L. (2002). Salvage esophagectomy for recurrent tumors after definitive chemotherapy and radiotherapy. *The Journal of Thoracic and Cardiovascular Surgery*, 123(1):175–183.
- Taketa, T., Correa, A. M., Suzuki, A., Blum, M. A., Chien, P., Lee, J. H., Welsh, J., Lin, S. H., Maru, D. M., Erasmus, J. J., Bhutani, M. S., Weston, B., Rice, D. C., Vaporciyan, A. A., Hofstetter, W. L., Swisher, S. G., and Ajani, J. A. (2012). Outcome of trimodality-eligible esophagogastric cancer patients who declined surgery after preoperative chemoradiation. *Oncology*, 83(5):300–4.
- Taketa, T., Sudo, K., Correa, A. M., Wadhwa, R., Shiozaki, H., Elimova, E., Campagna, M.-C., Blum, M. A., Skinner, H. D., Komaki, R. U., Lee, J. H., Bhutani, M. S., Weston, B. R., Rice, D. C., Swisher, S. G., Maru, D. M., Hofstetter, W. L., and Ajani, J. A. (2014). Post-chemoradiation surgical pathology stage can customize the surveillance strategy in patients with esophageal adenocarcinoma. *Journal of the National Comprehensive Cancer Network: JNCCN*, 12(8):1139–1144.
- Taketa, T., Xiao, L., Sudo, K., Suzuki, A., Wadhwa, R., Blum, M. A., Lee, J. H., Weston, B., Bhutani, M. S., Skinner, H., Komaki, R., Maru, D. M., Rice, D. C., Swisher, S. G., Hofstetter, W. L., and Ajani, J. A. (2013). Propensity-based matching between esophagogastric cancer patients who had surgery and who declined surgery after preoperative chemoradiation. *Oncology*, 85(2):95–99.
- Taylor, F. G. M., Quirke, P., Heald, R. J., Moran, B. J., Blomqvist, L., Swift, I. R., Sebag-Montefiore, D., Tekkis, P., Brown, G., and Magnetic Resonance Imaging in Rectal Cancer European Equivalence Study Study Group (2014).

- Preoperative magnetic resonance imaging assessment of circumferential resection margin predicts disease-free survival and local recurrence: 5-year follow-up results of the MERCURY study. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 32(1):34–43.
- Terheggen, G., Horn, E. M., Vieth, M., Gabbert, H., Enderle, M., Neugebauer, A., Schumacher, B., and Neuhaus, H. (2017). A randomised trial of endoscopic submucosal dissection versus endoscopic mucosal resection for early Barrett’s neoplasia. *Gut*, 66(5):783–793.
- Thomas, T., Singh, R., and Ragunath, K. (2009). Trimodal imaging-assisted endoscopic mucosal resection of early Barrett’s neoplasia. *Surgical Endoscopy*, 23(7):1609–1613.
- Vakil, N., Morris, A. I., Marcon, N., Segalin, A., Peracchia, A., Bethge, N., Zucaro, G., Bosco, J. J., and Jones, W. F. (2001). A prospective, randomized, controlled trial of covered expandable metal stents in the palliation of malignant esophageal obstruction at the gastroesophageal junction. *The American Journal of Gastroenterology*, 96(6):1791–1796.
- van der Valk, M. J. M., Marijnen, C. A. M., van Etten, B., Dijkstra, E. A., Hilling, D. E., Kranenbarg, E. M.-K., Putter, H., Roodvoets, A. G. H., Bahadoer, R. R., Fokstuen, T., Ten Tije, A. J., Capdevila, J., Hendriks, M. P., Edhemovic, I., Cervantes, A. M. R., de Groot, D. J. A., Nilsson, P. J., Glimelius, B., van de Velde, C. J. H., Hospers, G. A. P., and Collaborative investigators (2020). Compliance and tolerability of short-course radiotherapy followed by preoperative chemotherapy and surgery for high-risk rectal cancer - Results of the international randomized RAPIDO-trial. *Radiotherapy and Oncology: Journal of the European Society for Therapeutic Radiology and Oncology*, 147:75–83.
- van Gijn, W., Marijnen, C. A. M., Nagtegaal, I. D., Kranenbarg, E. M.-K., Putter, H., Wiggers, T., Rutten, H. J. T., Páhlman, L., Glimelius, B., van de Velde, C. J. H., and Dutch Colorectal Cancer Group (2011). Preoperative radiotherapy combined with total mesorectal excision for resectable rectal cancer: 12-year follow-up of the multicentre, randomised controlled TME trial. *The Lancet. Oncology*, 12(6):575–582.
- van Hagen, P., Hulshof, M. C., van Lanschot, J. J., Steyerberg, E. W., van Berge Henegouwen, M. I., Wijnhoven, B. P., Richel, D. J., Nieuwenhuijzen, G. A., Hospers, G. A., Bonenkamp, J. J., Cuesta, M. A., Blaisse, R. J., Busch, O. R., ten Kate, F. J., Creemers, G. J., Punt, C. J., Plukker, J. T., Verheul, H. M., Spillenaar Bilgen, E. J., van Dekken, H., van der Sangen, M. J., Rozema, T., Biermann, K., Beukema, J. C., Piet, A. H., van Rij, C. M., Reinders, J. G., Tilanus, H. W., van der Gaast, A., and Group, C. (2012). Preoperative chemoradiotherapy for esophageal or junctional cancer. *N Engl J Med*, 366(22):2074–84.

- van Vliet, E. P. M., Heijenbrok-Kal, M. H., Hunink, M. G. M., Kuipers, E. J., and Siersema, P. D. (2008). Staging investigations for oesophageal cancer: A meta-analysis. *British Journal of Cancer*, 98(3):547–557.
- Visbal, A. L., Allen, M. S., Miller, D. L., Deschamps, C., Trastek, V. F., and Pairolero, P. C. (2001). Ivor Lewis esophagogastrectomy for esophageal cancer. *The Annals of Thoracic Surgery*, 71(6):1803–1808.
- Vogel, J. D., Felder, S. I., Bhama, A. R., Hawkins, A. T., Langenfeld, S. J., Shaffer, V. O., Thorsen, A. J., Weiser, M. R., Chang, G. J., Lightner, A. L., Feingold, D. L., and Paquette, I. M. (2022). The American Society of Colon and Rectal Surgeons Clinical Practice Guidelines for the Management of Colon Cancer. *Diseases of the Colon and Rectum*, 65(2):148–177.
- Worrell, S. G., Alicuben, E. T., Oh, D. S., Hagen, J. A., and DeMeester, S. R. (2018). Accuracy of Clinical Staging and Outcome With Primary Resection for Local-Regionally Limited Esophageal Adenocarcinoma. *Annals of Surgery*, 267(3):484–488.
- Wouters, M. W. J. M., Karim-Kos, H. E., le Cessie, S., Wijnhoven, B. P. L., Stassen, L. P. S., Steup, W. H., Tilanus, H. W., and Tollenaar, R. a. E. M. (2009). Centralization of esophageal cancer surgery: Does it improve clinical outcome? *Annals of Surgical Oncology*, 16(7):1789–1798.
- Yang, H., Liu, H., Chen, Y., Zhu, C., Fang, W., Yu, Z., Mao, W., Xiang, J., Han, Y., Chen, Z., Yang, H., Wang, J., Pang, Q., Zheng, X., Yang, H., Li, T., Lordick, F., D’Journo, X. B., Cerfolio, R. J., Korst, R. J., Novoa, N. M., Swanson, S. J., Brunelli, A., Ismail, M., Fernando, H. C., Zhang, X., Li, Q., Wang, G., Chen, B., Mao, T., Kong, M., Guo, X., Lin, T., Liu, M., Fu, J., and AME Thoracic Surgery Collaborative Group (2018). Neoadjuvant Chemoradiotherapy Followed by Surgery Versus Surgery Alone for Locally Advanced Squamous Cell Carcinoma of the Esophagus (NEOCRTEC5010): A Phase III Multicenter, Randomized, Open-Label Clinical Trial. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 36(27):2796–2803.
- Ychou, M., Boige, V., Pignon, J.-P., Conroy, T., Bouché, O., Lebreton, G., Ducourtieux, M., Bedenne, L., Fabre, J.-M., Saint-Aubert, B., Genève, J., Lasser, P., and Rougier, P. (2011). Perioperative chemotherapy compared with surgery alone for resectable gastroesophageal adenocarcinoma: An FNCLCC and FFCD multicenter phase III trial. *Journal of Clinical Oncology: Official Journal of the American Society of Clinical Oncology*, 29(13):1715–1721.
- Zhang, J. Q., Hooker, C. M., Brock, M. V., Shin, J., Lee, S., How, R., Franco, N., Prevas, H., Hulbert, A., and Yang, S. C. (2012). Neoadjuvant chemoradiation therapy is beneficial for clinical stage T2 N0 esophageal cancer patients due to inaccurate preoperative staging. *Ann Thorac Surg*, 93(2):429–35; discussion 436–7.

- Zhao, X., Ren, Y., Hu, Y., Cui, N., Wang, X., and Cui, Y. (2018). Neoadjuvant chemotherapy versus neoadjuvant chemoradiotherapy for cancer of the esophagus or the gastroesophageal junction: A meta-analysis based on clinical trials. *PloS One*, 13(8):e0202185.
- Zhou, C., Zhang, L., Wang, H., Ma, X., Shi, B., Chen, W., He, J., Wang, K., Liu, P., and Ren, Y. (2015). Superiority of Minimally Invasive Oesophagectomy in Reducing In-Hospital Mortality of Patients with Resectable Oesophageal Cancer: A Meta-Analysis. *PloS One*, 10(7):e0132889.