



# Principles of surgical oncology, reconstructive surgery and oncological rehabilitation.



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# Hungarian Task Force Against Cancer

**Content** (WHO-Recommendation):

- Primer prevention
- Secunder prevention
- Early diagnostic
- Therapy
- Rehabilitation
- Palliation
- Education
- Public Relation
- Attendance
- Structure of Oncological Providers
- Monitoring, Quality Measurement

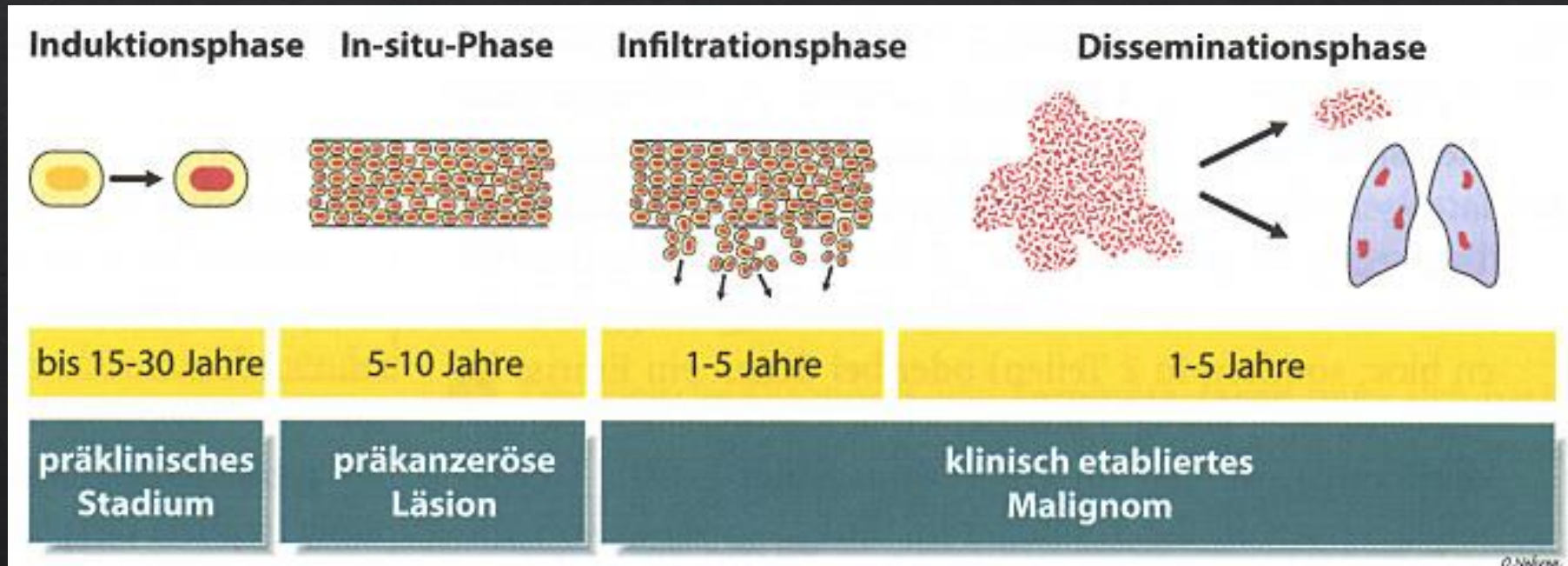
# Principles of tumors

- ◇ Originated from cells of the body
- ◇ Presence and growth of tumors have no advantage for the individuum
- ◇ Growth of them are independent from normal controlling systems
- ◇ Without therapy this condition leads into death

# Etiology

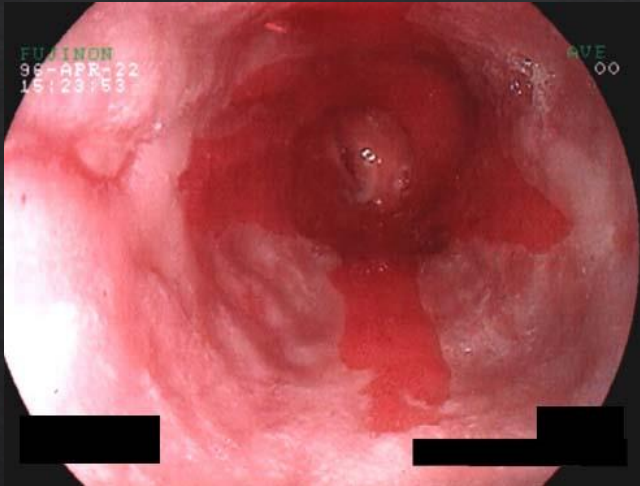
- ◇ Physical conditions
- ◇ Chemical agents
- ◇ Viruses (for example: HCV, HPV)
- ◇ Precancerous conditions
- ◇ Genetical factors

# 4-stage idea for tumor growth



# Precancerous condition

**Conditions that (may) lead to cancer growth**



**Barrett-  
metaplasia**

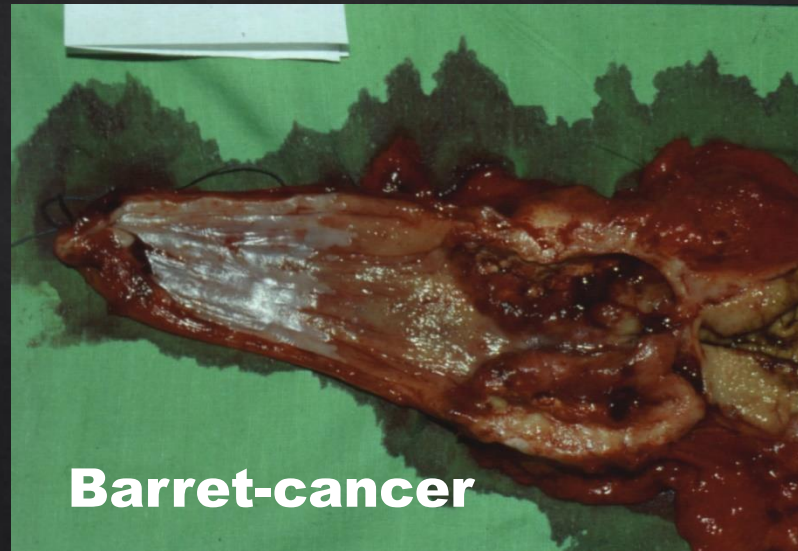
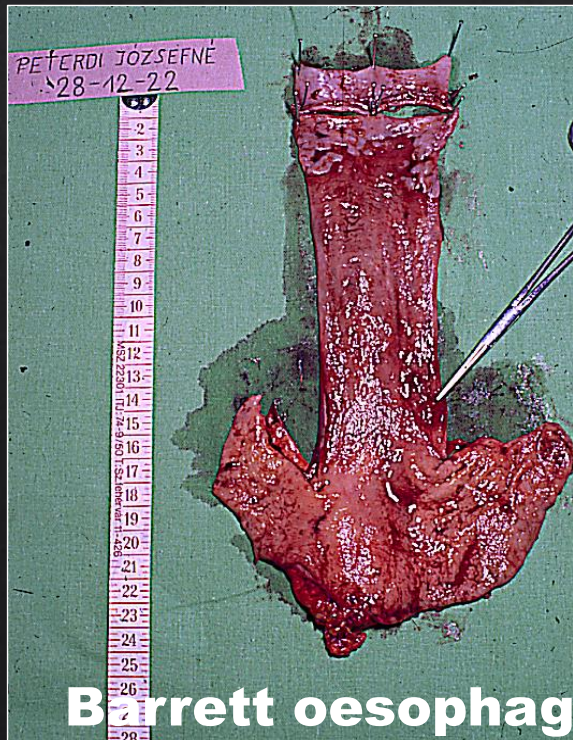
**Polyposus coli**



**Cervicalis  
Intraepithelialis  
neoplasia (CIN)**

# Precancerous laesion

**Histologically verified condition, which may lead with a higher probabilty (or even sure way) to malignant tumor.**



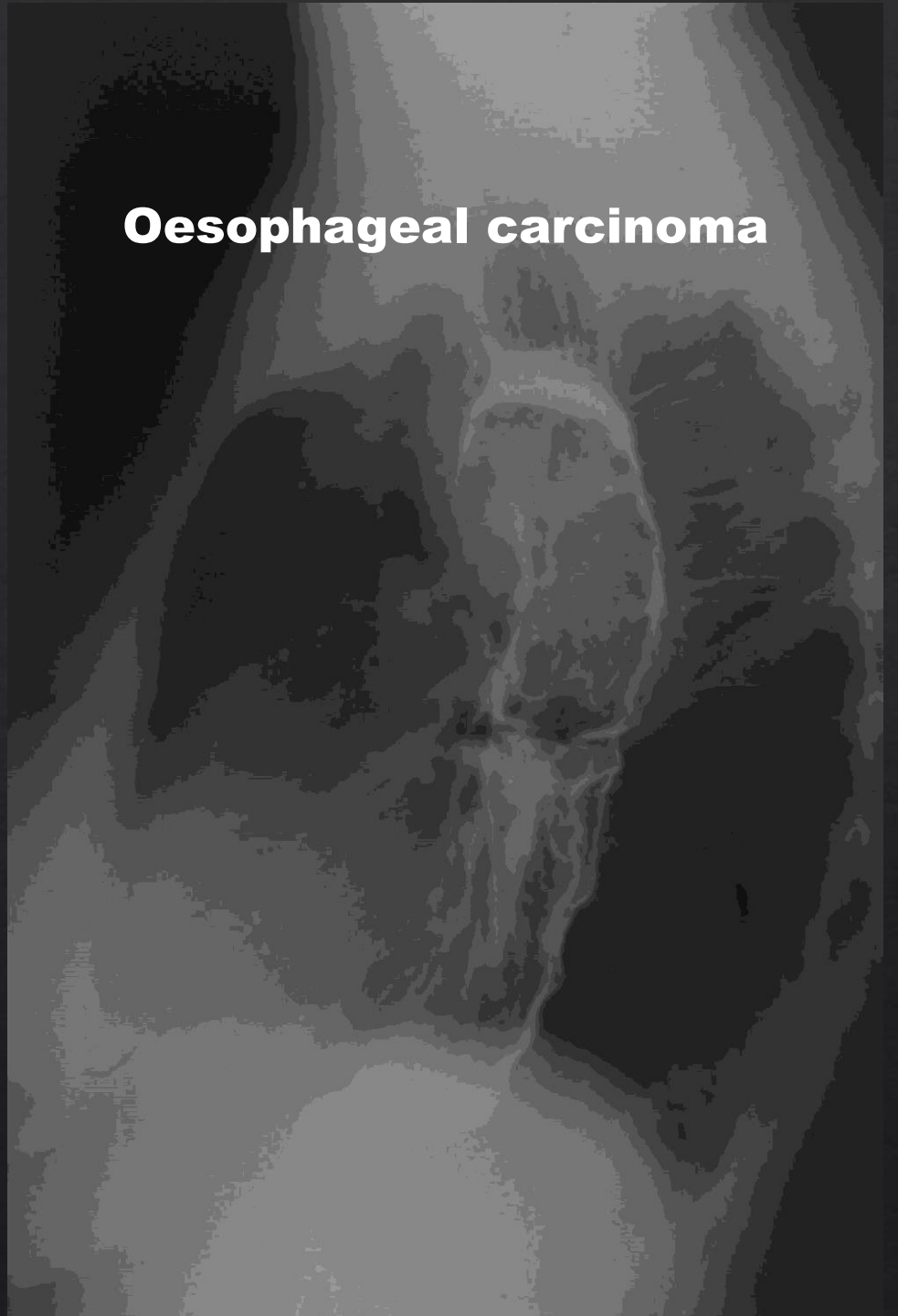
**Familial colon polyposis**

*Barium swallow*

**Gastric ulcer**



**Oesophageal carcinoma**





Anal carcinoma (HPV)



# What are the questions of the clinicians?

Is there a malignancy in the body? What is the histology?

**Staging?**

1. Where is the tumor situated?
2. What are the borders of the tumor?
3. Is it infiltrating the neighbourhood?
4. Is there local or distant metastasis?

**Grading?**

1. Which grade of differentiation the tumor have?
2. Grade of atypia?
3. Mitotic rate per window in histological report?

**Other prognostic factors?**

1. Free margin?
2. Invasion into blood or lymphatic vessels?
3. What is the rate of spreading?
4. Vasculra pattern of tumors?
5. Proliferations markers?
6. Immunhisztochemical prognostic factors?

**Tumorbiological conditions?**

1. Is it a hereditary tumor?
2. Genetical abnormalities?
3. Predictive factors?
4. Resistency for therapy? (hormonstatus, receptorstatus)
5. Metabolic rate of the tumor?
6. Oxigenisation of the tumor?
7. Viral factors influencing the tumor?

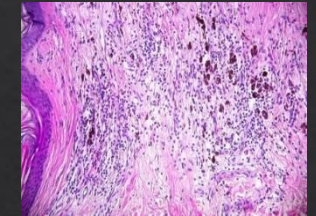
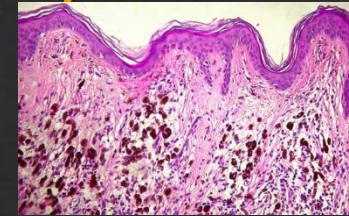
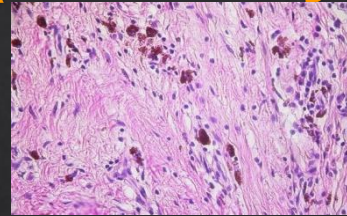
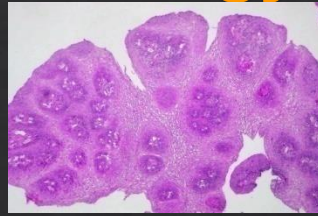
**Relaps?**

1. Staging?
2. Grading?
3. Tumorbiology?

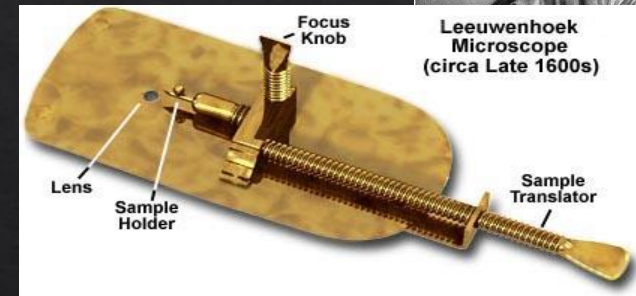
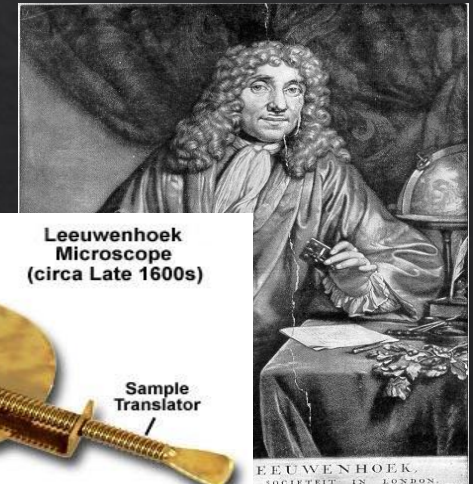
# Pathology (classic type)

## Morphology

- ◇ Cytology
- ◇ Core biopsy
- ◇ Intraoperativ frozen section (resection margins and lymph node involvement)
- ◇ Hystological verification (morphology and immunohystochemical fenotype)
- ◇ Specific genetical abnormalities from morphology?



- Pathological staging (pT, pN)
  - Tumor size
  - Presence of metastases in amount and place
- Patological grading
  - Differentiation
  - Atypia of the nucleus
  - Mitotic activity
- Other pathological prognostic factors
  - Integrity and size of free margin
  - Infiltration pattern of tumors: Expansiv – infiltrativ; vessel, lymphatic vessel, perineural growth
  - Proliferation markers: Ki67 proliferation index, DNS cytometry
  - Immunhystochemical prognostic factors: ER, PR status, Tp53 pozitivness



# Molecular pathology

- ◇ Support pathological report
  - ◇ Genetical analysis (lymphomas, sarcomas)
  - ◇ Viral DNA proof
- ◇ Predictive factors
  - ◇ Ploidia (DNS cytometry, kromosoma number)
  - ◇ Clonal examinations (lymphomas)
  - ◇ Génmutations (hereditary tumors)
- ◇ Prognostic factors
  - ◇ Geneexpression
    - ◇ Tissue microarray, mRNS microarray
  - ◇ Geneamplification (for example: breast cancer, neuroblastoma)
  - ◇ Microsatellita instability (pl. CRC)
  - ◇ Onkogene mutations (pl. EGFR, KRAS, BRAF)
- ◇ Targeted anticancer therapy
  - ◇ Breast cancer, lung cancer, colorectal cancer, GIST, lymphoma, melanoma malignum, etc.

# Staging

1. Physical exam: relevant exams
2. Anatomical imaging (static or dynamic informations)
  - X-ray
  - angiography (DSA)
  - multilayer digital technologies
    - US
    - CT-MDCT
    - MRI (1,5T>)
3. Pathological staging
  - section: tumor size and stage
  - hystology: presence of malignoma
  - molecular path exams (verifications of diagnosis)
    - viral verification
    - genetic pattern (lymphomák, szarkómák)
4. Functional imaging (molecular imaging)
  - metabolic activity
  - izotop diagnostic tools (SPECT, PET, PET-CT)
  - MRSI, dyn MR, DMR, perf. CT/MR, organ specific. KA-MR

# Grading

1. Rate of differentiation: - histology

2. Rate of atypia: - histology

## 3. **Pathologic prognostic factors**

1. Free margin: - histology

2. Vascular invasion: - histology

3. Vascular pattern, angiogenesis:  
- histology

4. - Immunohistochemistry

- endothel marker

- proliferation markers: index, DNS cytometry (PET-CT, MRSI)

- dyn CT/MR, perfusion CT/MR, USD, Ka-US

5. Growth speed and pattern: - histology

6. Immunohistochemical prognostic factors:

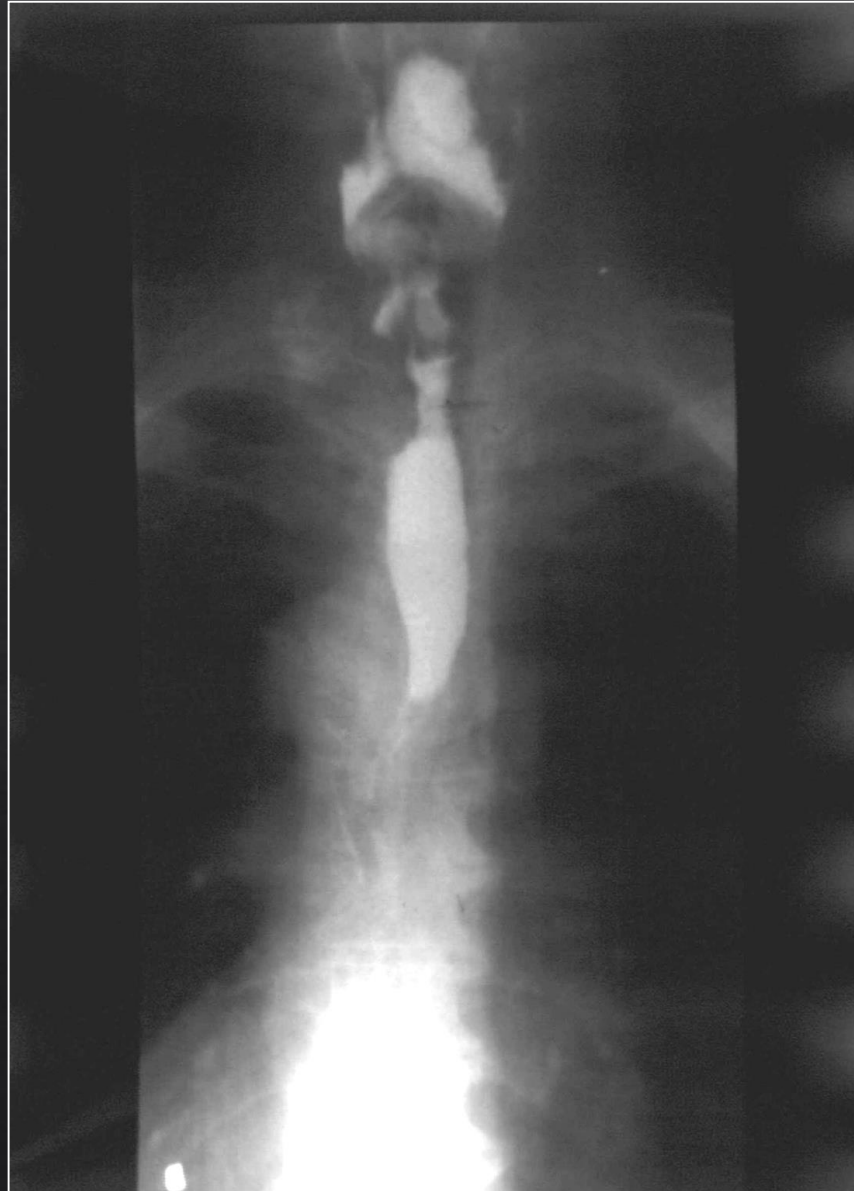
- ER, PR status, Tp 53 positiveness

# Tumor specific symptoms

- ◇ Changing in habity of stool or urinate
- ◇ Non-healing wound
- ◇ Unusual bleeding or growth
- ◇ Node formations or growth
- ◇ Dysphagy
- ◇ Lymph node enlargement
- ◇ Cough

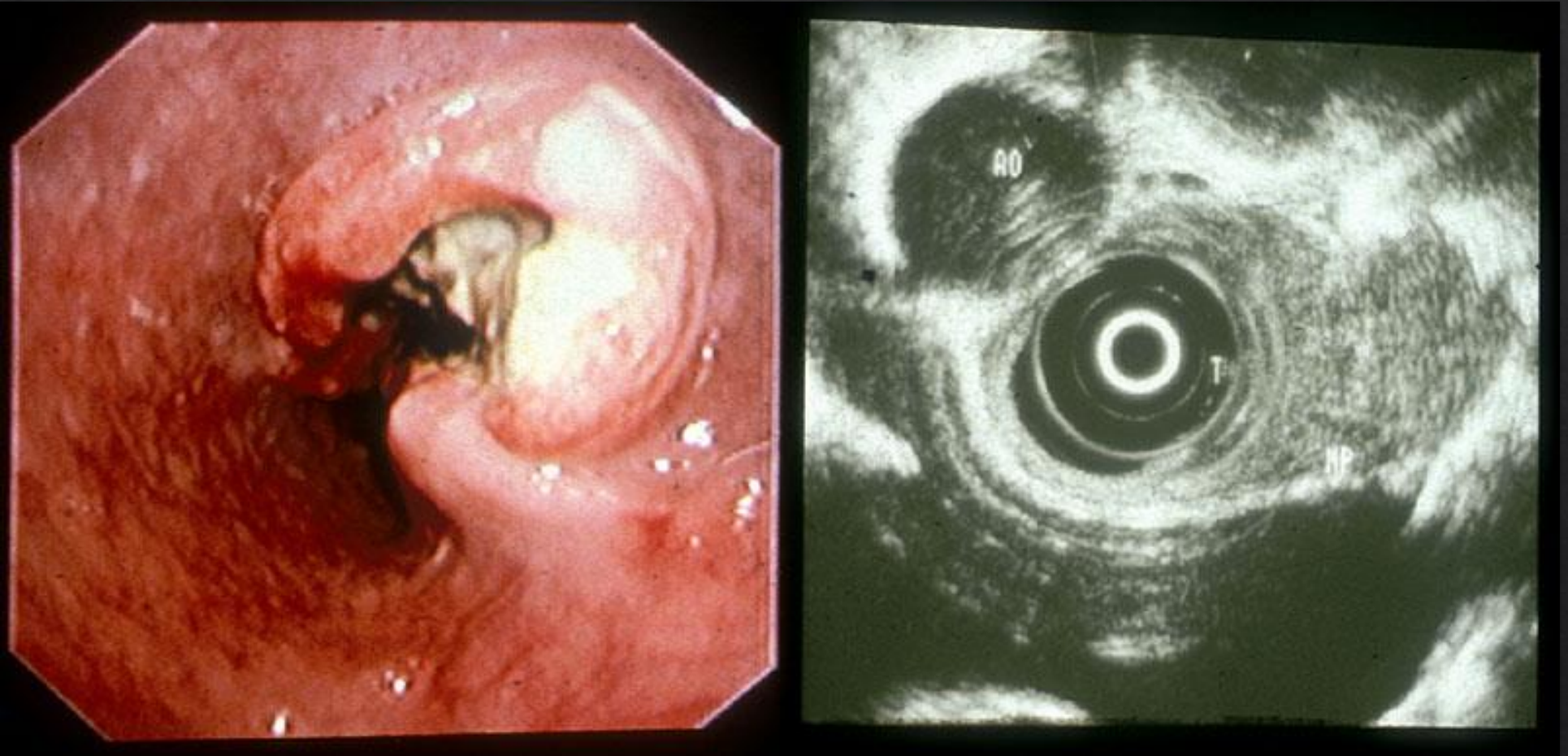
# Radiology

Oesophageal  
carcinoma  
(upper third)



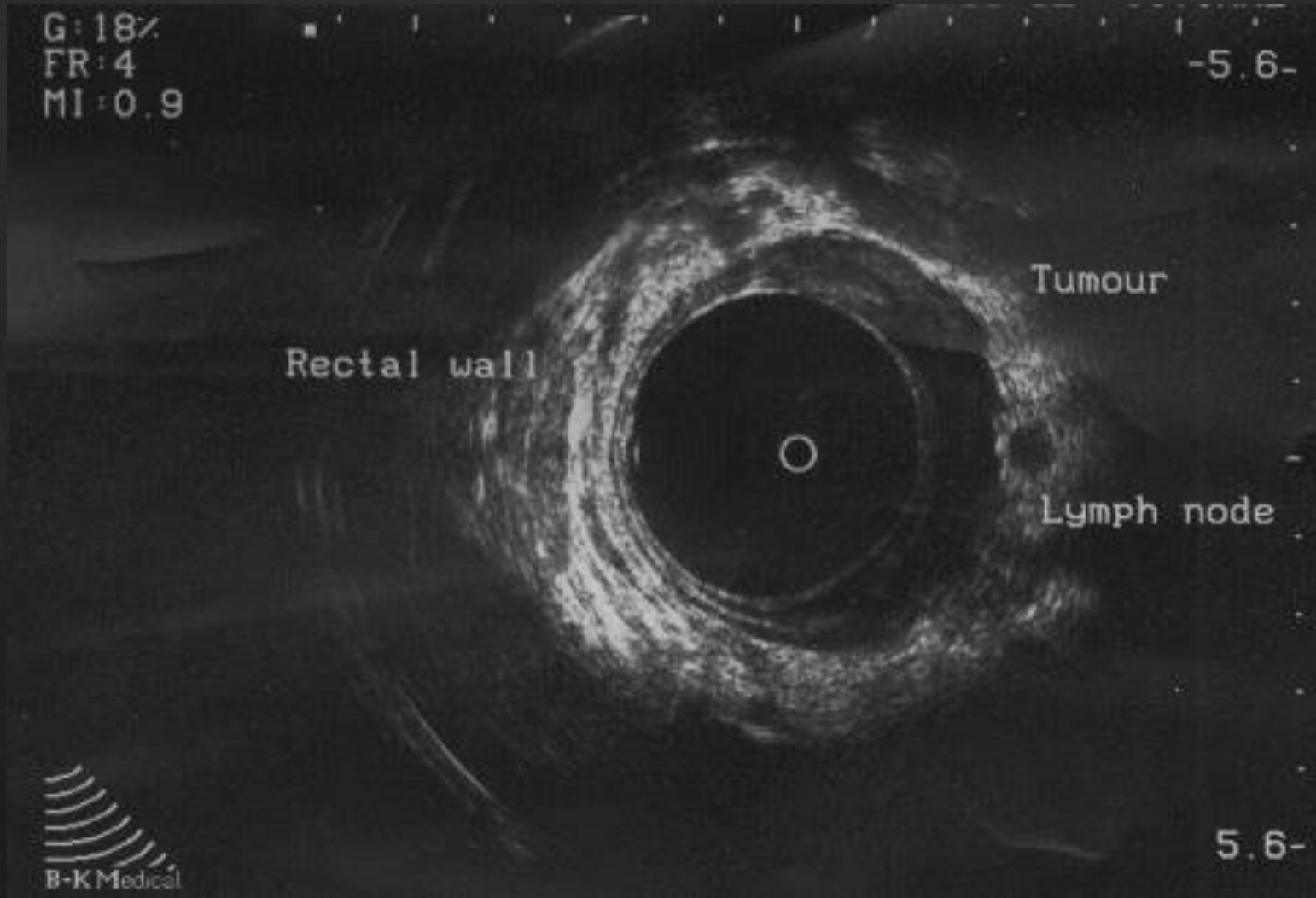


# Endoscopy and endoscopic US



**Oesophageal carcinoma**

# Endorectal US



ORSOS JANOSNE  
HV21/3/51  
21-MAR-1951  
13:30:18  
15-FEB-1999  
IMAGE 13

A

Pecsi Diagnosztikai Kp.  
SOMATOM AR  
AB3 5 H-SP-CR VD10E

**CT**

**Liver tumor**



SCAN 13  
TI 1.3 s  
mA 100  
kV 110  
SL 10.0  
TP -166  
FoV 274  
CE 12/0  
GT 0

W 174  
C 32

NAGY FERENC KAROLYNE,,,,

ID: 067474017

\* 1954.02.15.

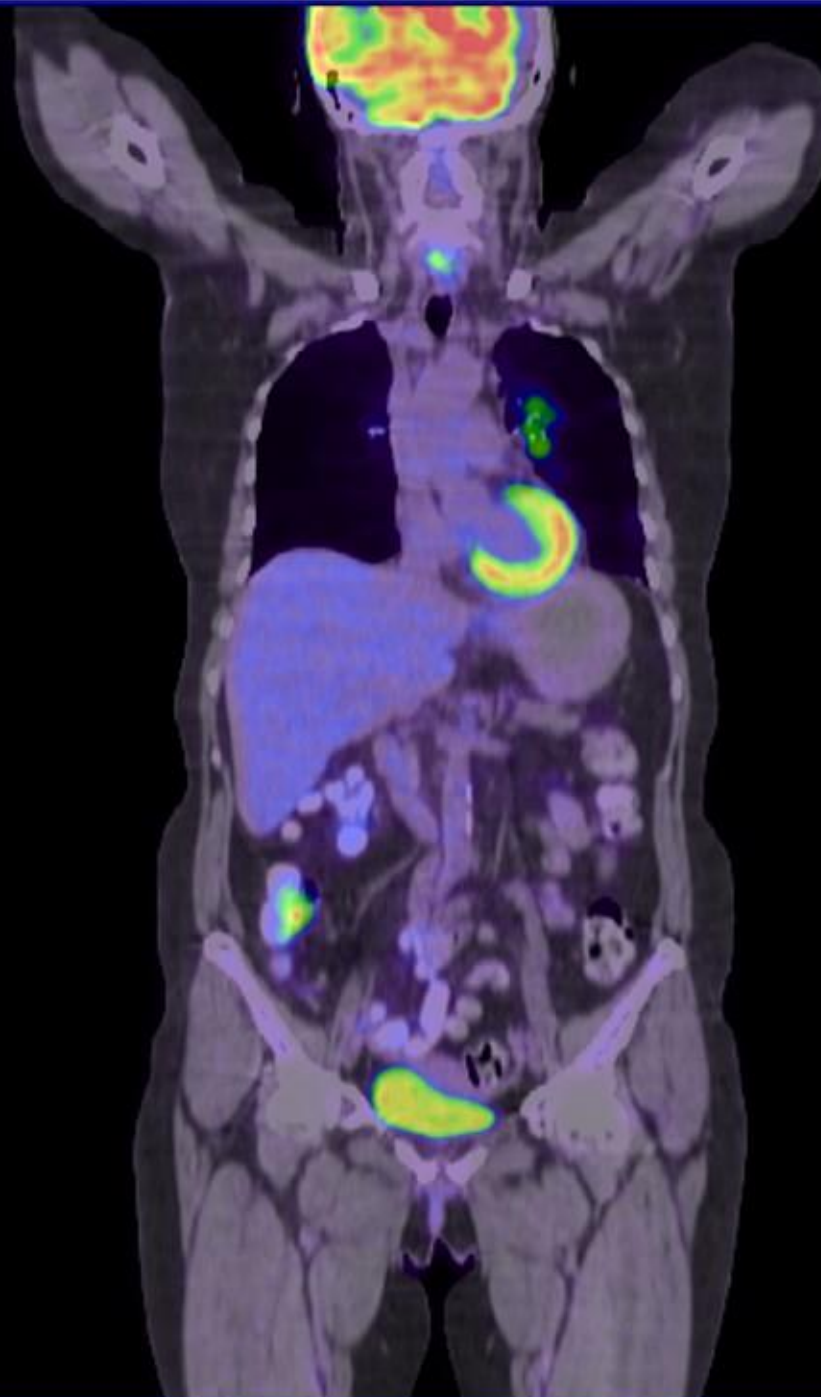
Study 1

2010.09.07.

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16 IMA

Pozitron Diagnosztika  
Biograph 6



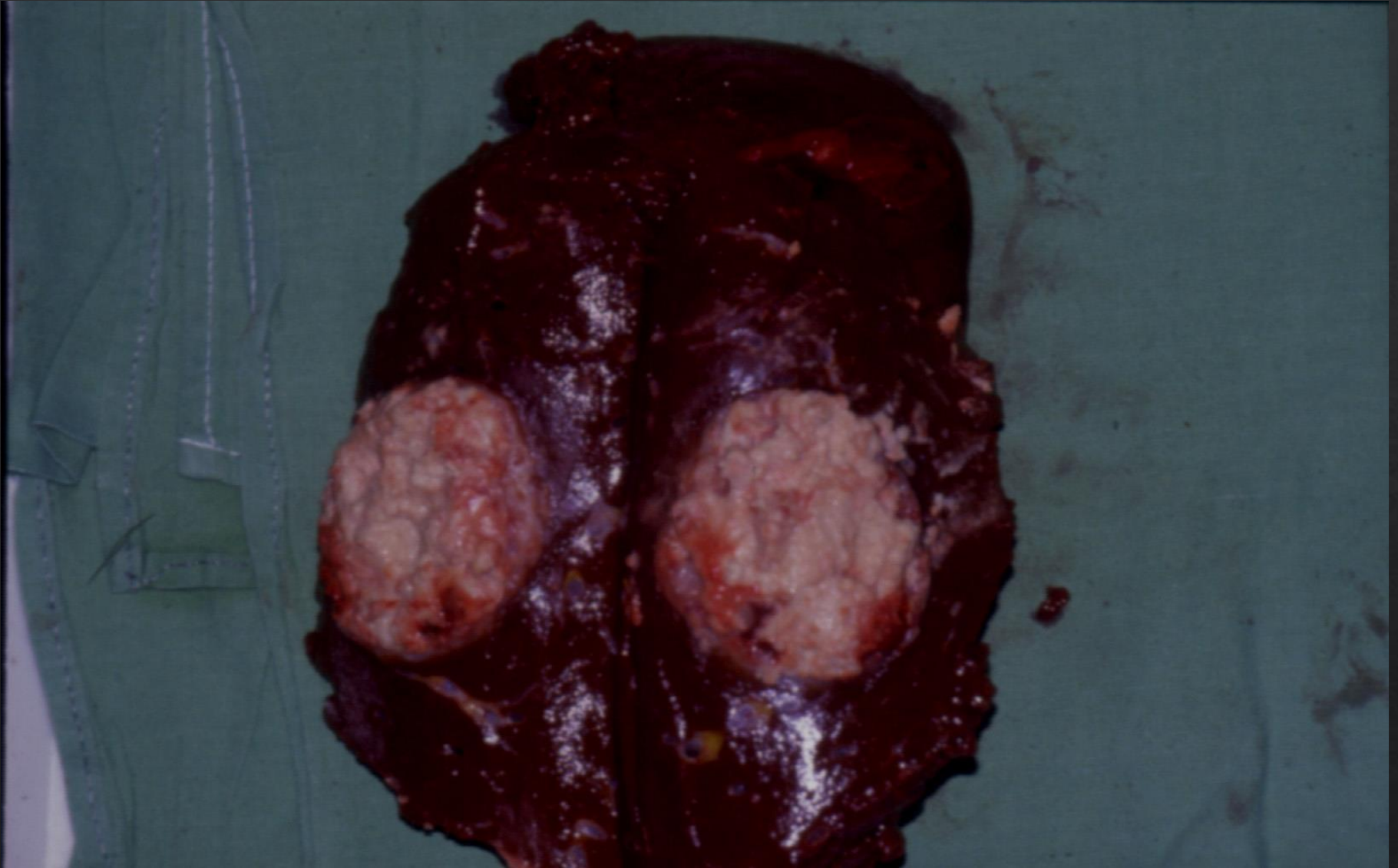
PET-CT



# TNM system

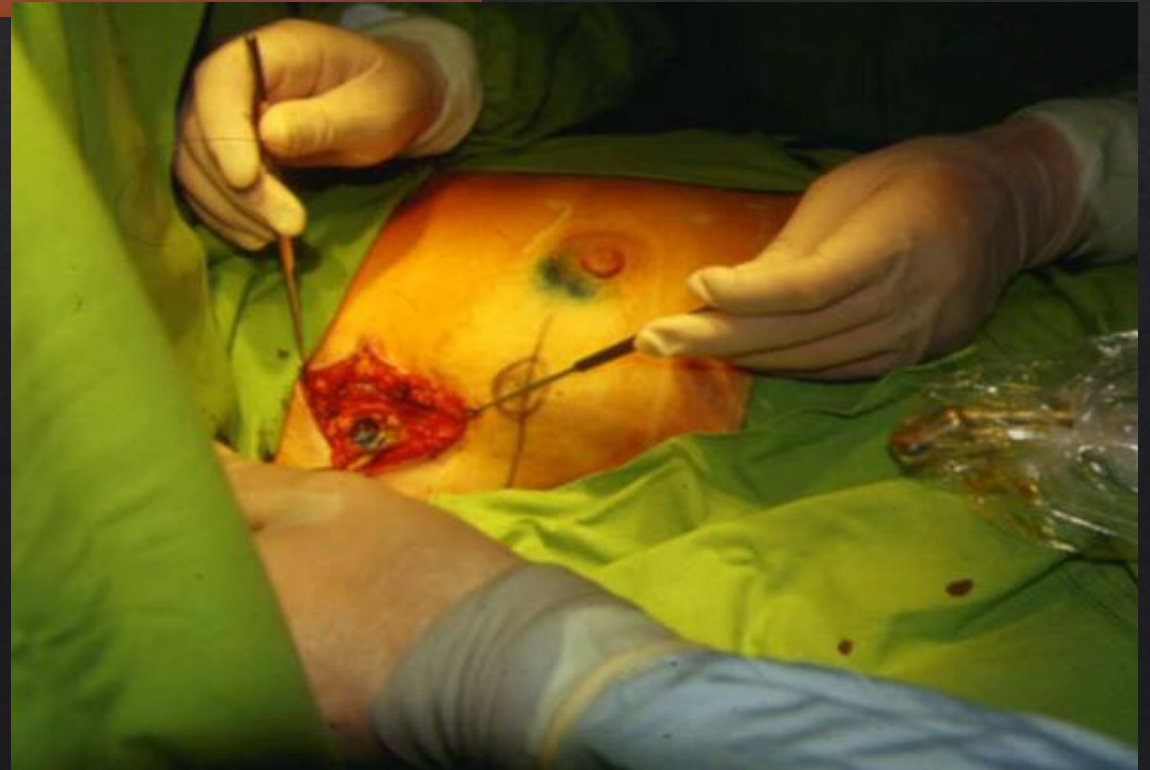
- T tumor size
- N is the regional lymph nodal involvement
- M is the distant metastases
  
- cTNM- clinical TNM (based on radiology)
- pTNM- pathological TNM (hystological proff)
- ypTNM –pathological TNM after therapy

# Liver met





Sentinel lymph node  
biopsy



# Multidisciplinary therapy

- ◇ **Surgery**
- ◇ Radiotherapy
- ◇ Systemic therapy
  - ◇ Chemotherapy, Hormonotherapy, Immunotherapy, targeted therapy
- ◇ Combined (multidisciplinary) therapy:
  - ◇ Surgery+radiotherapy (stomach, rectum)
  - ◇ Surgery and synchronous radiotherapy (breast)
  - ◇ Neoadjuvant therapy: Radio+/-chemo+ surgery (rectum, stomach, oesophagus)
  - ◇ Systemic treatment, surgery of metastasis, neoadjuvant radio-chemotherapy, surgery of the primary tumor, adjuvant chemotherapy (rectumcc+livermetastasis)



# MDT team

## Consist of:

- medical oncologist
- pathologist
- surgeon
- radiotherapist



# The role of MDT in oncological patient's care

## 1. Determining diagnostic algorithm

1.1. Evaluate of the existing reports

1.2. Complementation of existing reports

- Is there a therapeutic cosequency?
- What is the goal?
- Sensitivity and specificity of chosen examination?  
(CT, MRI, PET CT)
- Sequence of diagnosis?
- Where to do?

## 2. Determining therapeutic approaches

- Therapeutic forms to declare  
(surgery, irradiation, systemic)
- Sequence of the therapies
- Timing of therapies
- Place of therapies

# The role of MDT in oncological patient's care

## 3. Control of therapy

- What?
- How?
- When?
- Where?

## 4. Rehabilitation-palliation

### 4.1. Physical:

- pain management
- reconstructive surgery (breast, face, skin, extremity)
- type
- time
- stoma-care (trachea, GI tract, urogenital tract)
- lymphoedema care
- necrosis and fibrosis of skin or mucosa
- side-effect of chemotherapy

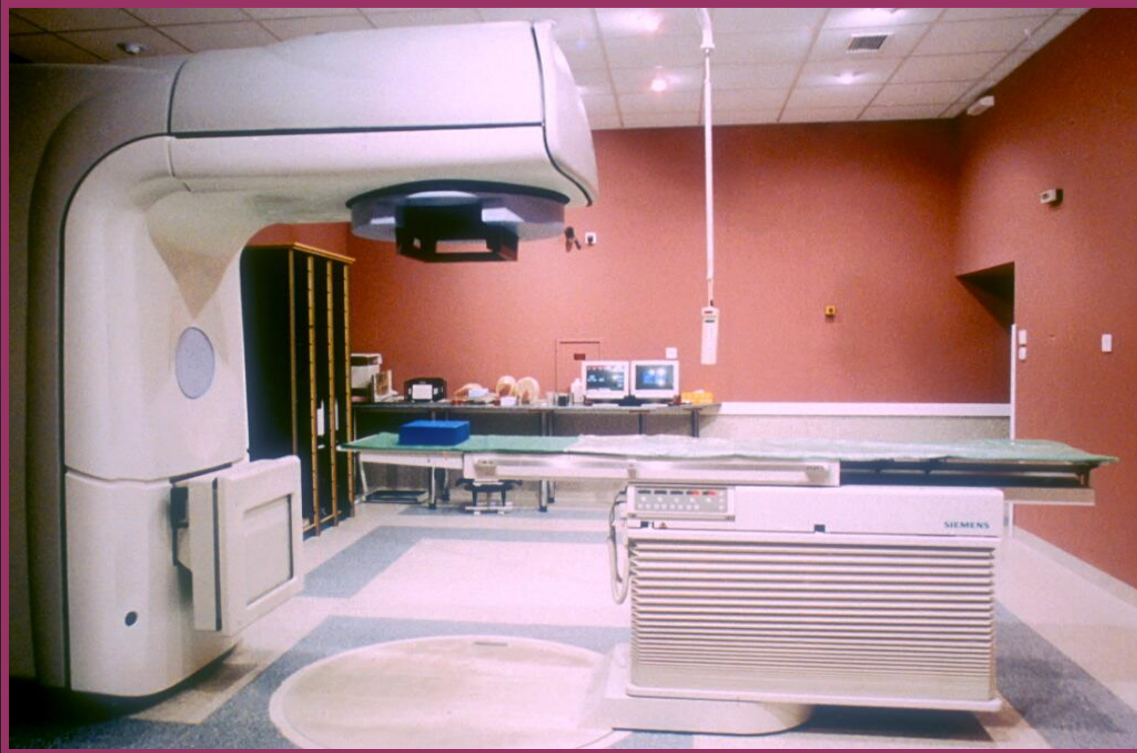
### 4.2. Emotional

## 5. Task:

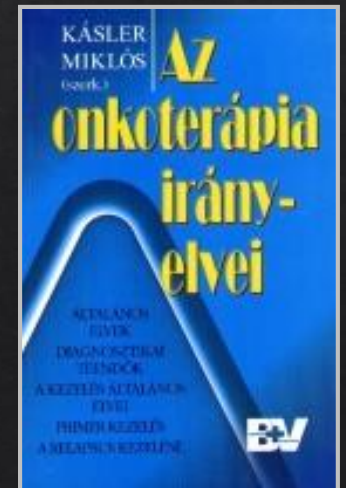
laws, permissions

- obligatory
- operational field

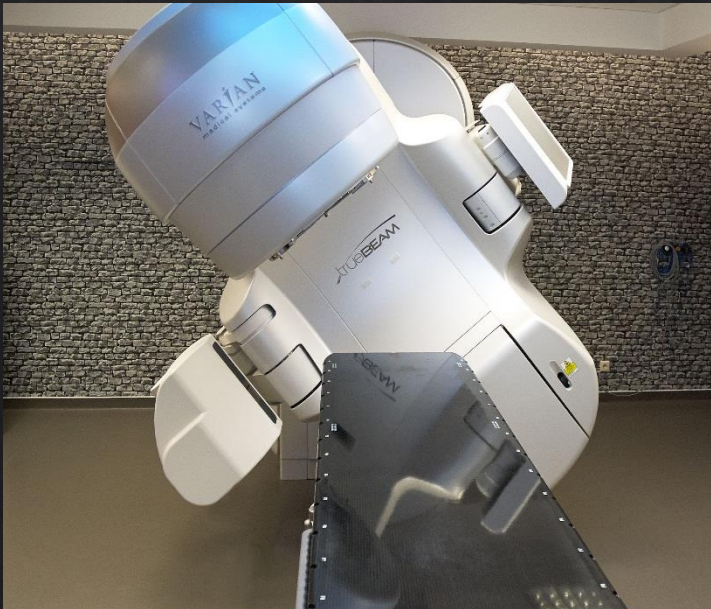
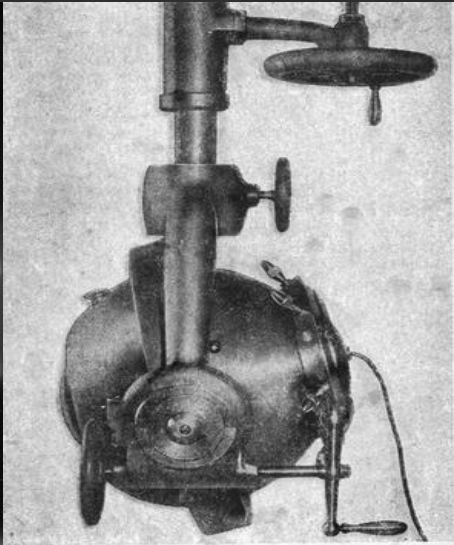
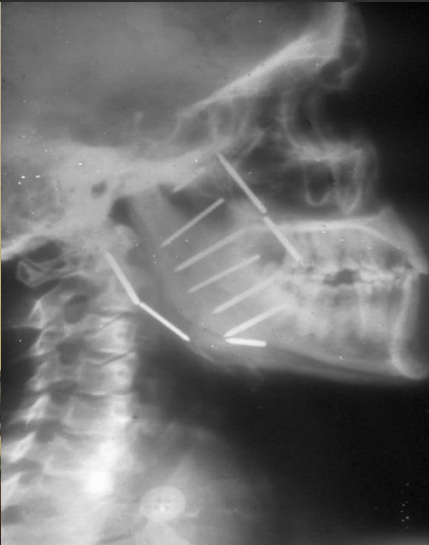
# KOMPLEX -CARE



1. Surgery
2. Irradiation
3. Chemotherapy

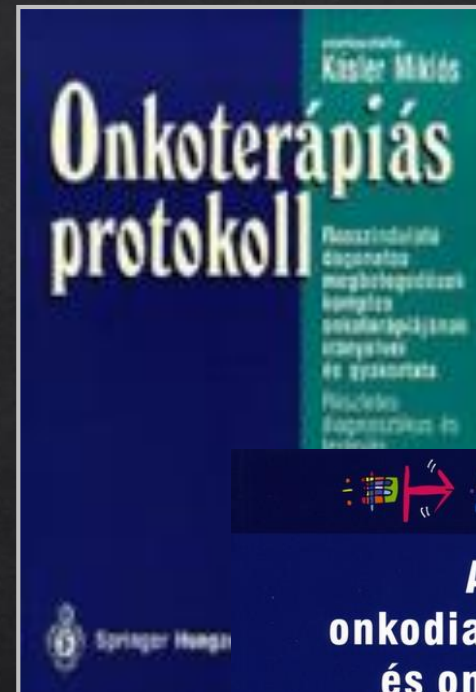


# Irradiation



# Chemo

1. **Cytostatic**
2. **Targeted**
3. **Immuno**





**Chemotherapy**

**Biological therapy**



# Oncosurgery

- ◇ 1. prophylactic
- ◇ 2. diagnostic
- ◇ 3. therapeutical
- ◇ 4. palliative

## Prophylaktic oncosurgery

- **1. Rectal polyps**
- **2. Hereditary bowel cancer**
- **3. Colitis ulcerosa**
- **4. Hereditary medullar thyroid cancer**
- **6. BRCA-positiv breast cancer**
- **7. Hereditary ovarial cancer**



# Surgery

## ◇ 1. Curative intent

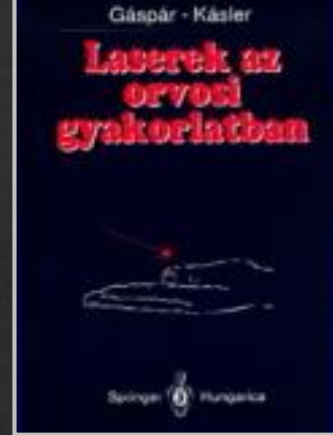
- ◇ Operability – For the patients
- ◇ Resecability – For the tumor
- ◇ Organ-sparing vs. radicality
- ◇ Free surgical margin is essential = R0
- ◇ Quality of life
- ◇ Reconstruction Immediate /delayed /Oncoplastic surgery
- ◇ Minimal Invasive Surgery (MIS) – Laparoscopia, SILS, NOTES, TEM, Robotic-surgery (da Vinci)

## ◇ 2. Palliation

- ◇ -stoma-formation, bypass (GEA, ileo-transversostomia), vessel-ligature etc.

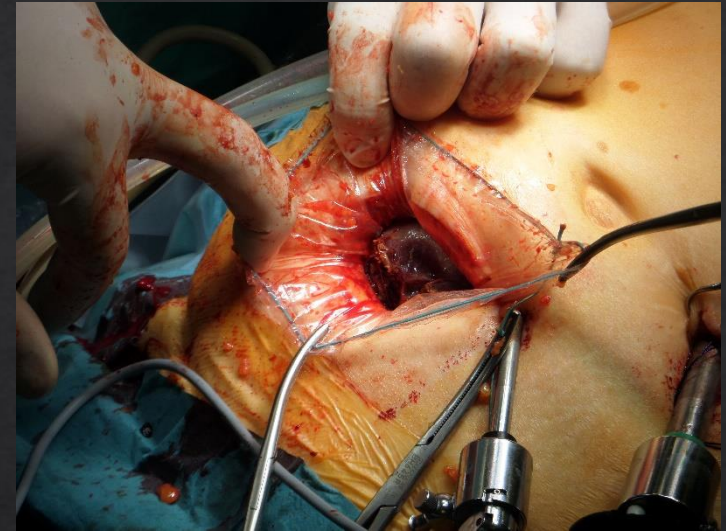
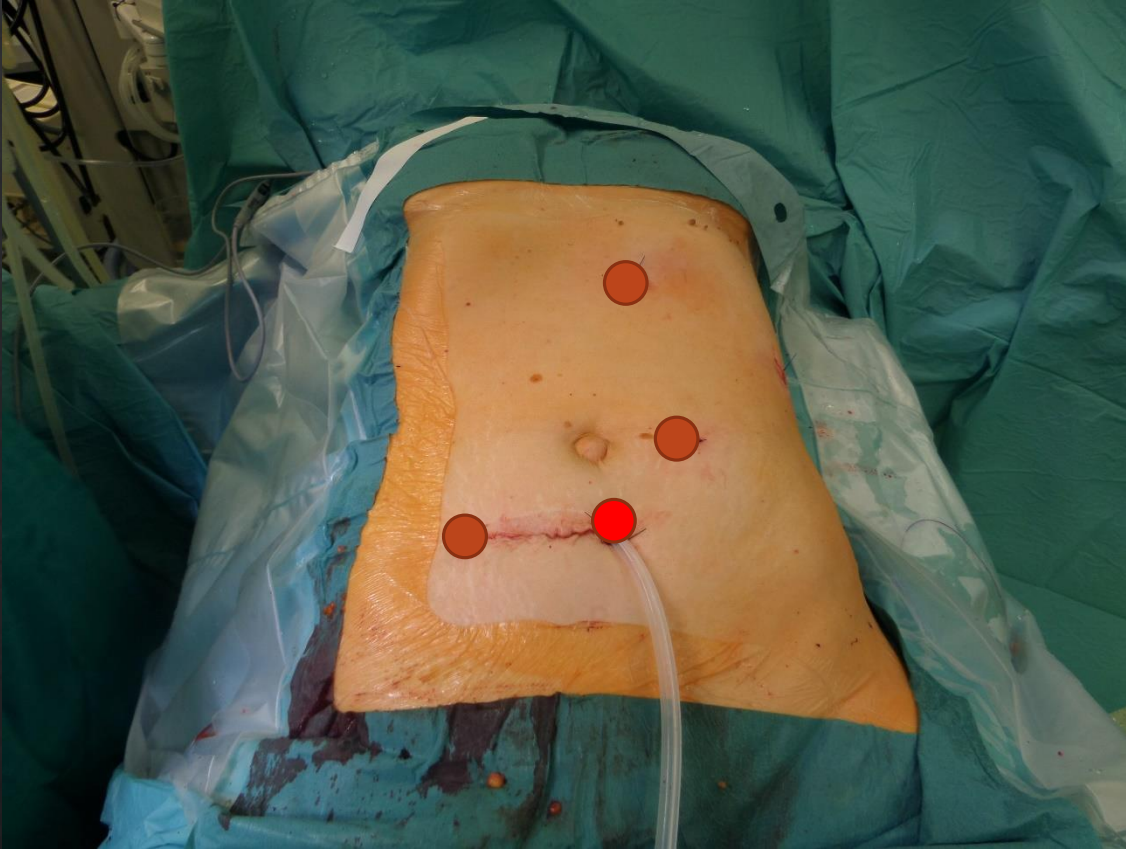
# Principles of oncological surgery

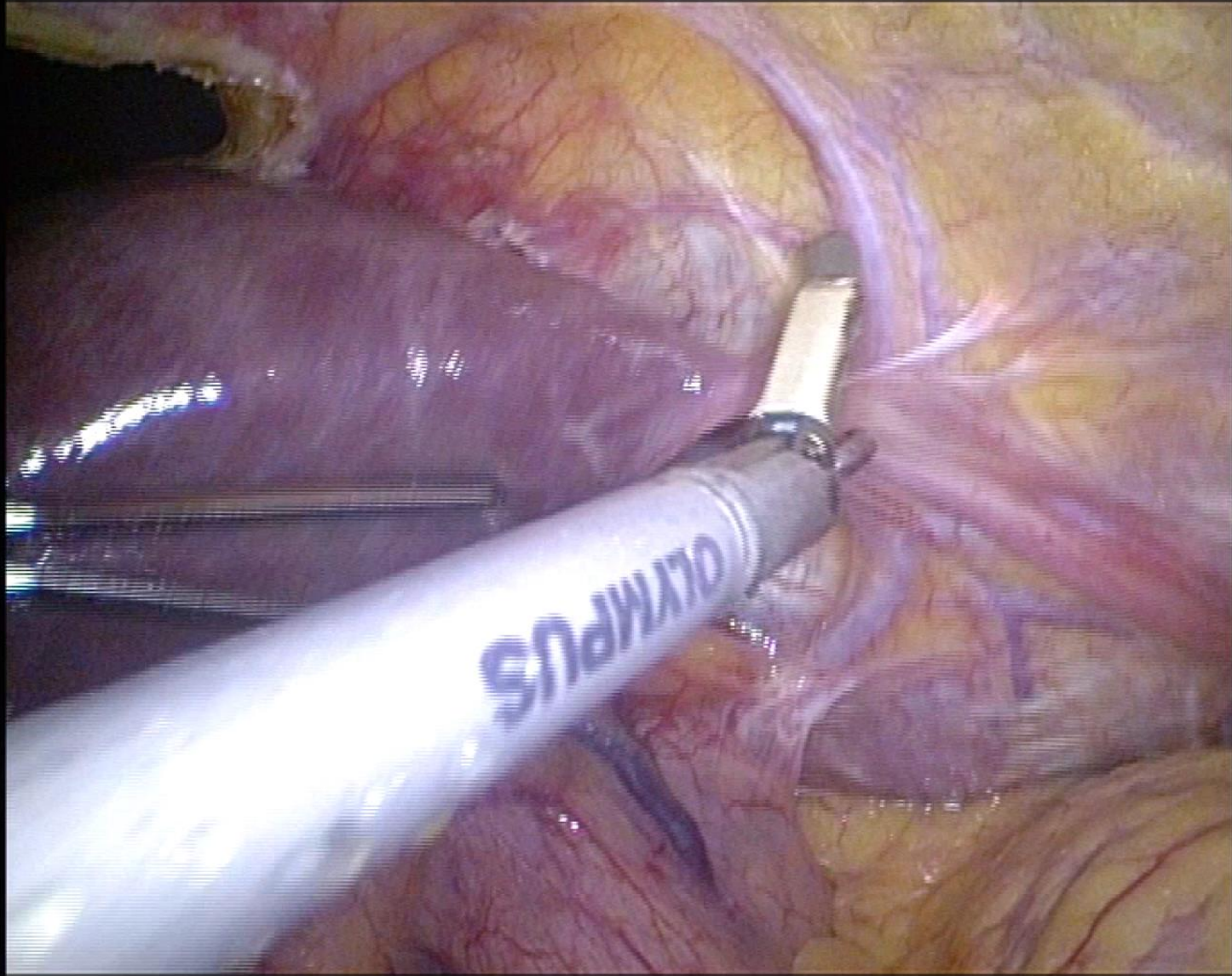
1. Radicality (“en block”)
  - 1.1. Primary Tumor
  - 1.2. Metastases
2. Monoblock Operation
  - 2.1. Lymphogen spreading
  - 2.2. Hematogen spreading
3. “No touch technique”
  - 3.1. Access to the operation field (minimal invasiv surgery)
  - 3.2. Operational technique
  - 3.3. Instrumental arsenal (Laser, high-energy sealer device)
4. Unity of structure-function and esthetic
5. Reconstruction of the structure-function (esthetic)



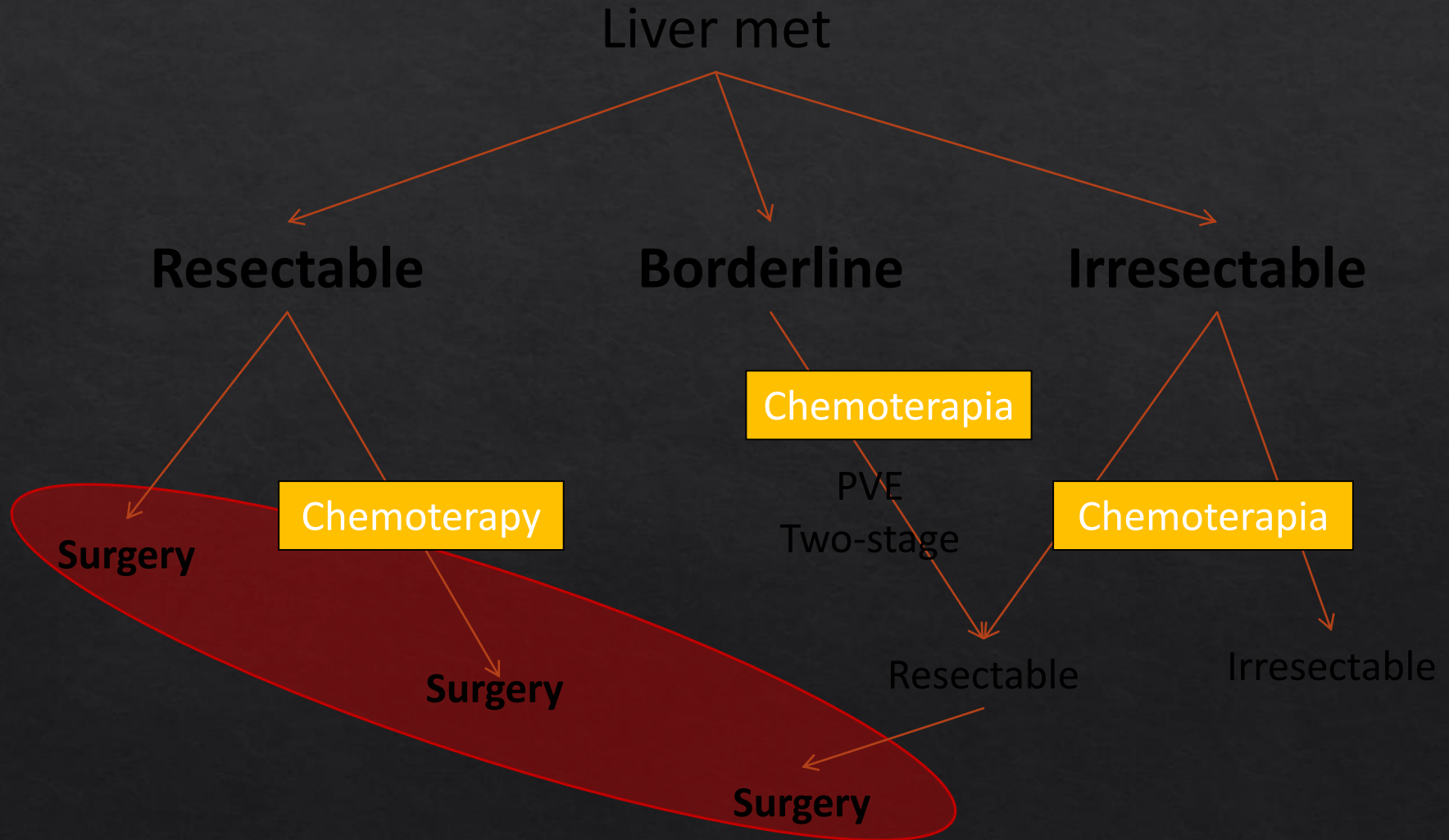


# Laparoscopic liver surgery





# Strategy (CRC liver metastasis)



# Evolution of oncosurgery I.

## 1. Primer Tumor

### 1.1. Decreased radicality

- Structure
- Function
- Estheticum

### 1.2. Komplex Terapy

- Radiation therapy
- Chemo

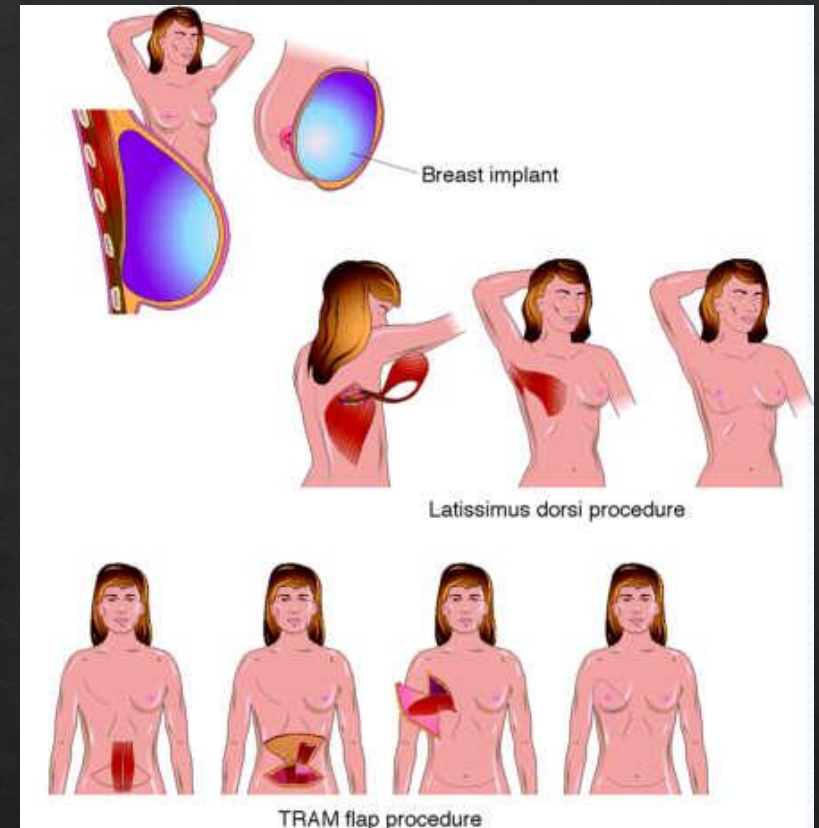
## 2. Metastases

### 2.1. Regional Metastasis

- Sentinel lymph node biopsy
- Block-dissection

### 2.2. Resection of mets

- Systemic control over the disease, metastases are resectable
- Met is soliter, or situated in one region
- „high risk” patients
- Mutlidisciplinary approach!



# Evaluation of oncosurgery II.

## 3. Reconstruction

### 3.1. Site

- 3.1.1. Head-neck region
- 3.1.2. Breast
- 3.1.3. Skin, neuronal, muscle
- 3.1.4. GI tract



### 3.2. Principles

- 3.2.1. Improvement of QoL (body, soul, function-esthetic)
- 3.2.2. Replacement with similar organs
- 3.2.3. Immediate reconstruction

### 3.3. Type of reconstruction

#### 3.3.1. Autologous transplantation

- flaps (neighbourhood or distant)
- Fasciocutan flap
- Myocutan flap
- Osteomyocutan flap
- Other visceral organ

#### 3.3.2. Heterogenous transplantation

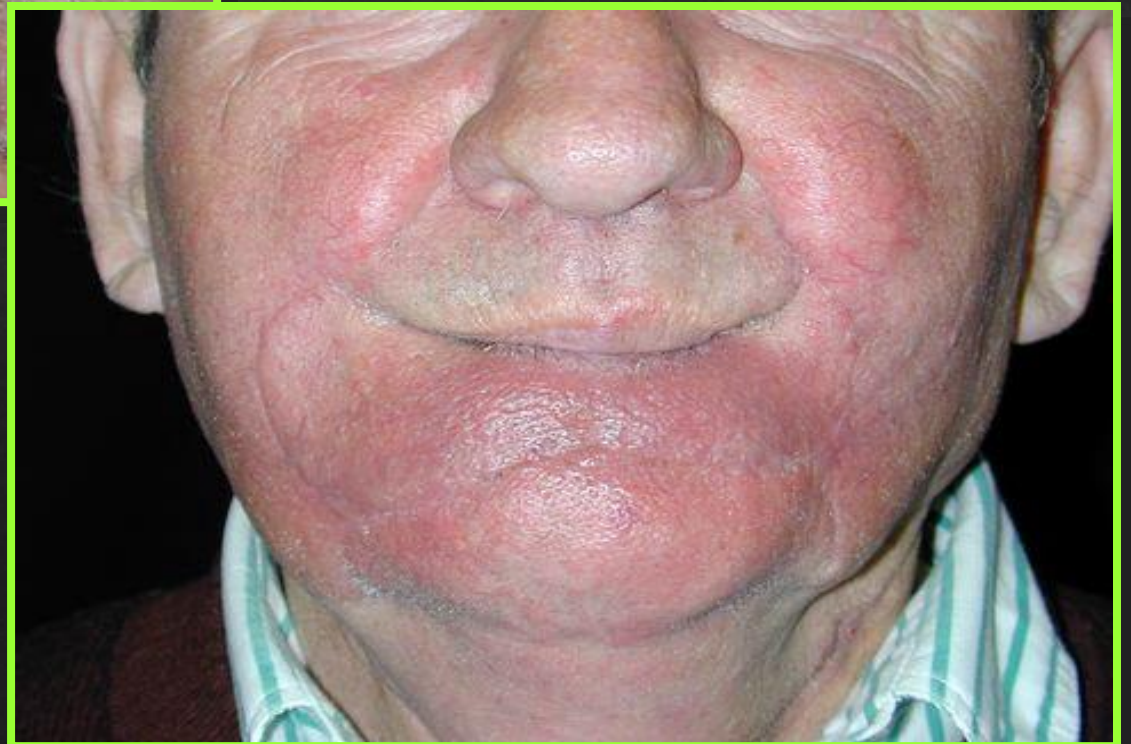
- Prothesis
- Expander





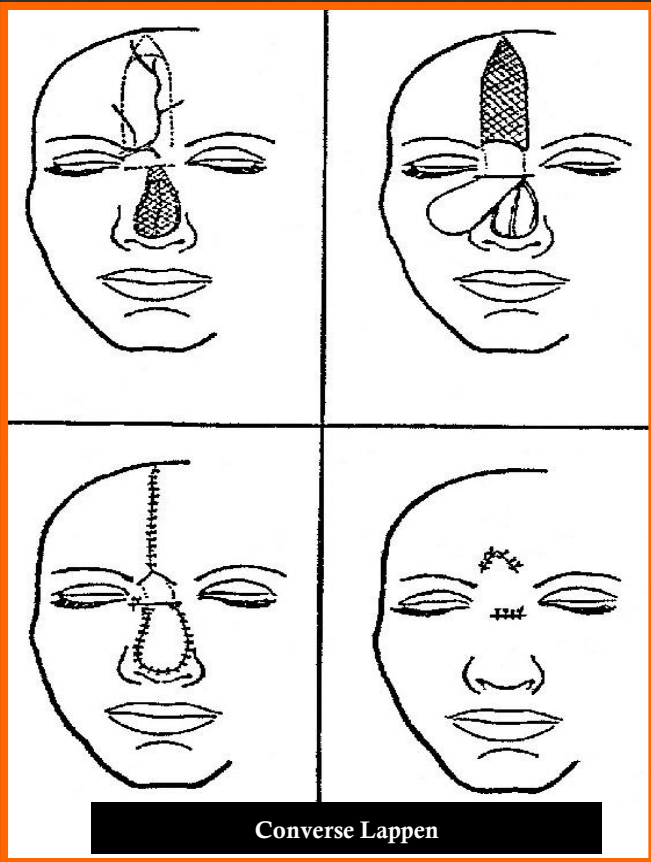


**Lower limb reconstruction  
with local flap**









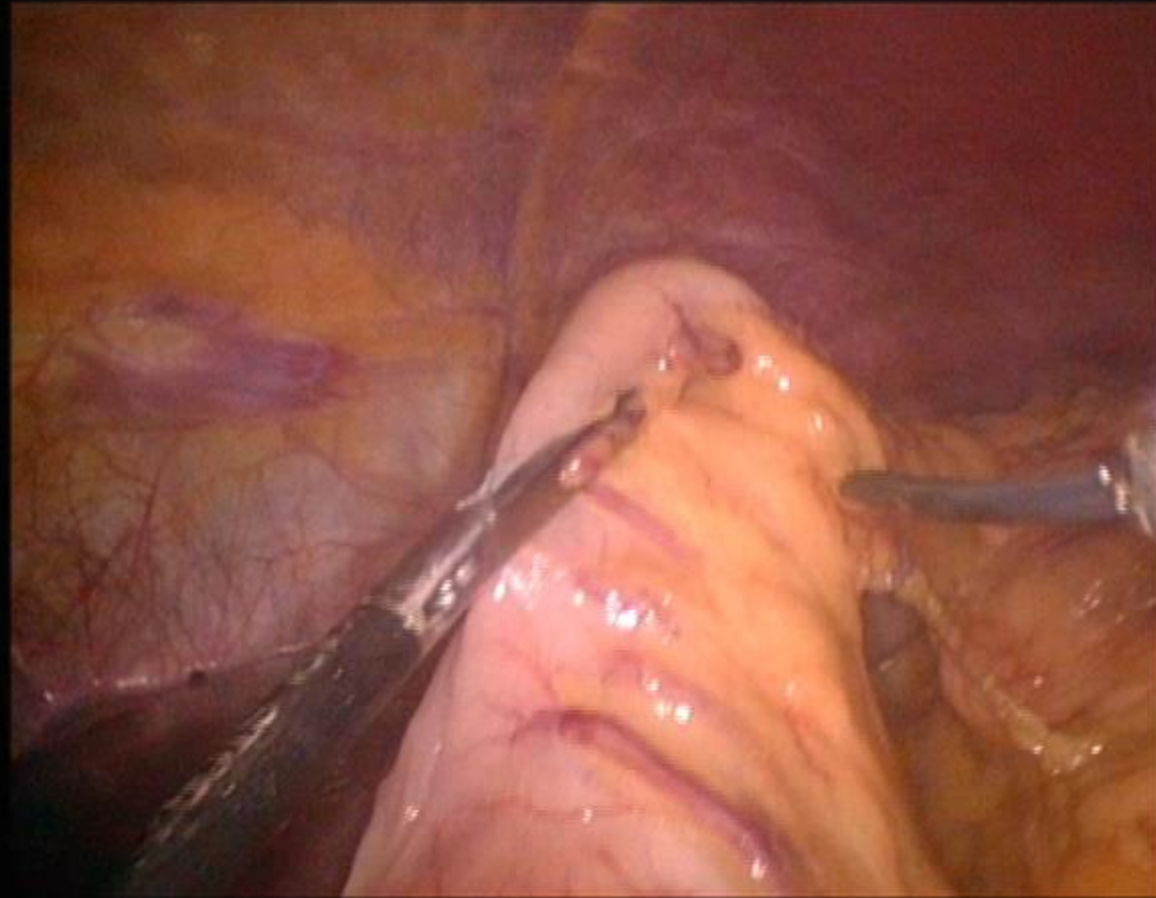
# Converse Flap



# MI Oesophagus (Thorax)



# Oesophagus replacement with stomach



# Free transplantation

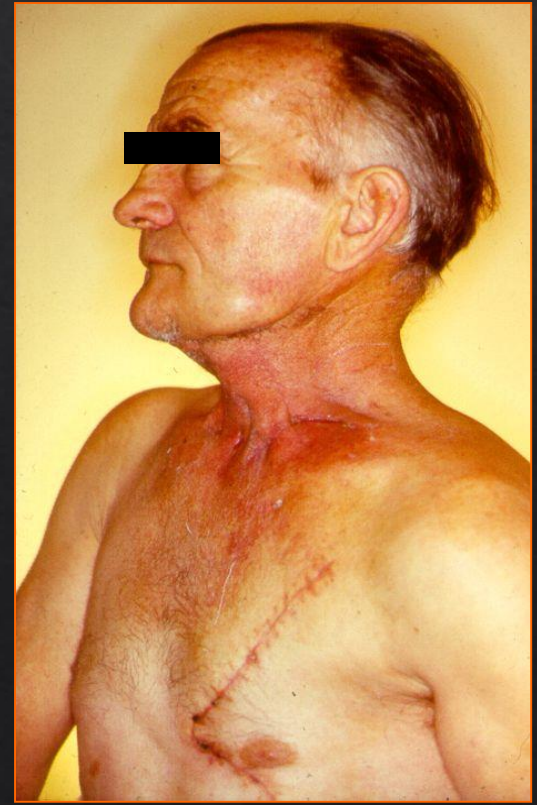
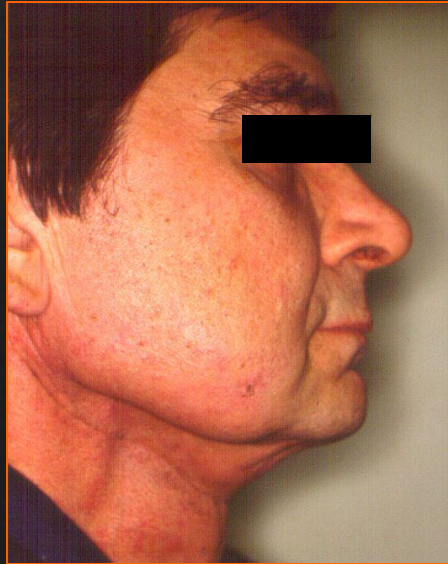
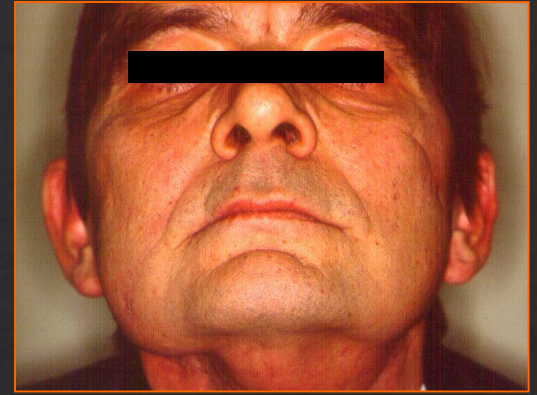
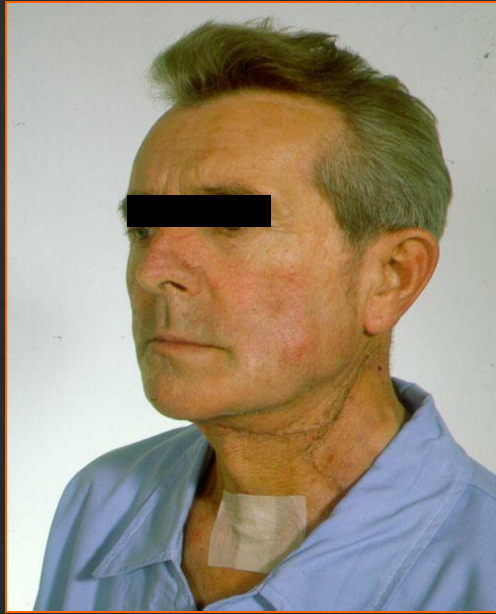
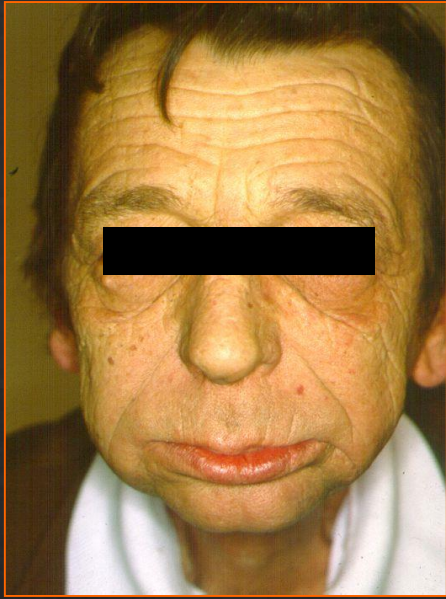
## Types:

- Skin
- Half-thick skin
- Free jejunal interpositum with microvascular anastomosis
- Myocutan (osteomyocutan, osteocutan) flap transplantation with microvascular anastomosis

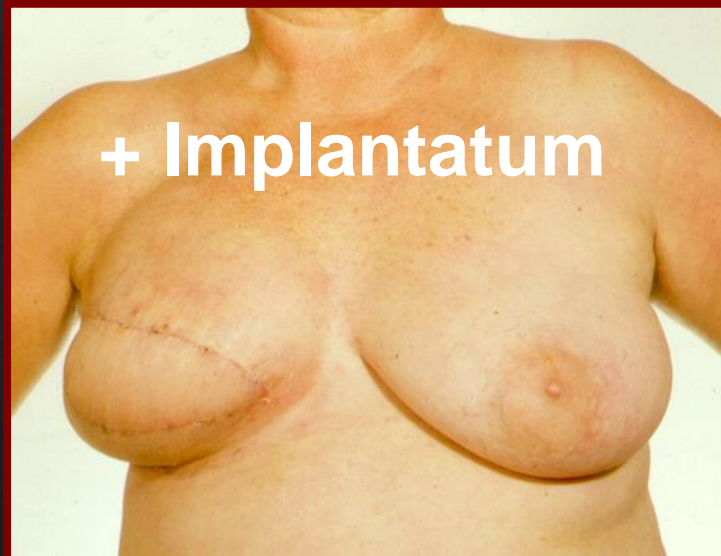


# Skintumor – Half-thick skin transposition

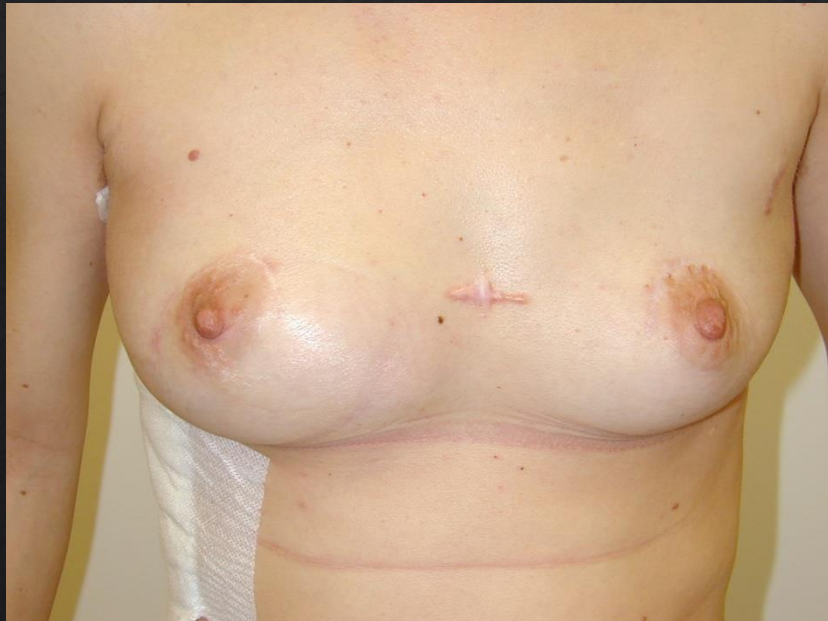


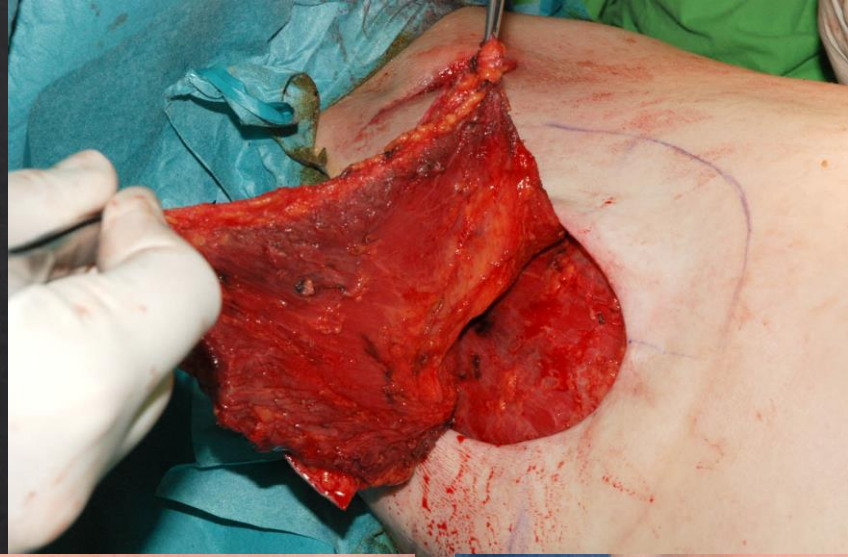


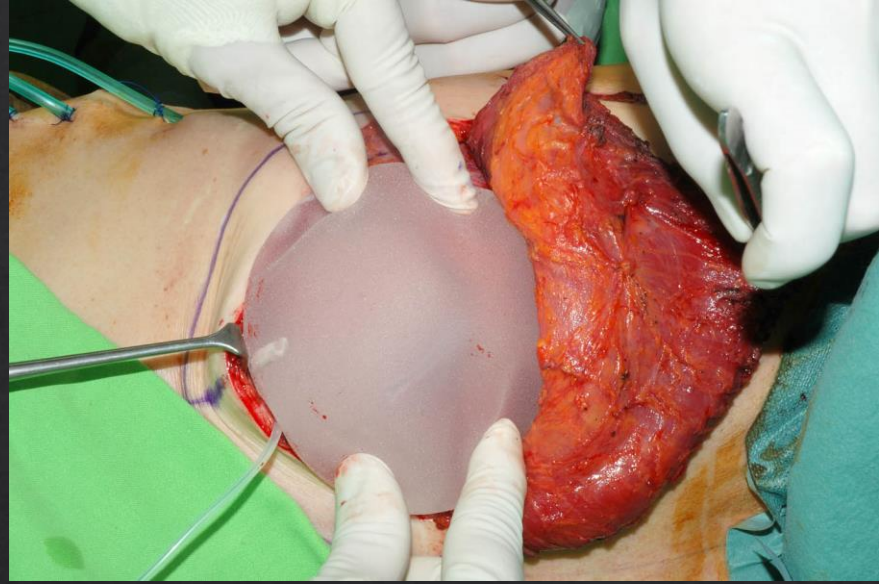
# Latissimus dorsi musculocutan flap



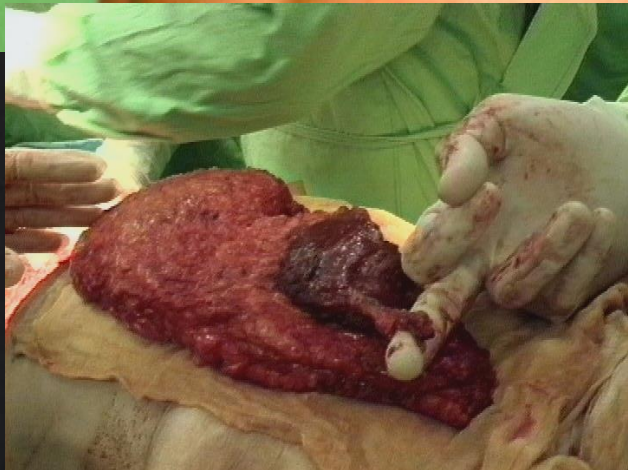
# Latissimus dorsi musculocutan flap



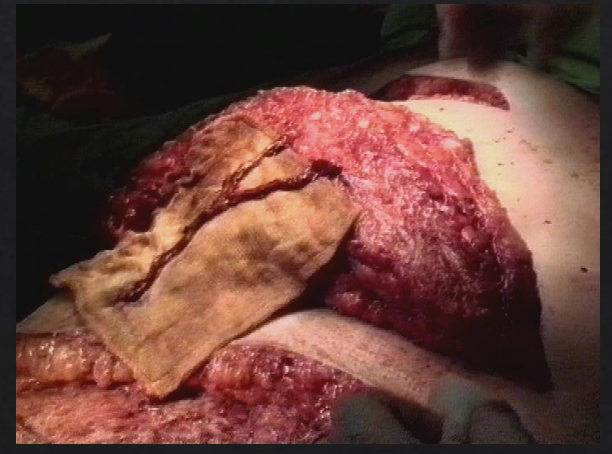
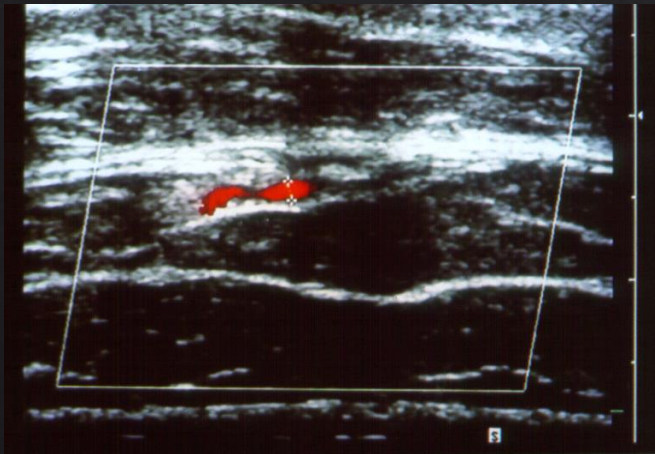




# Free TRAM flap

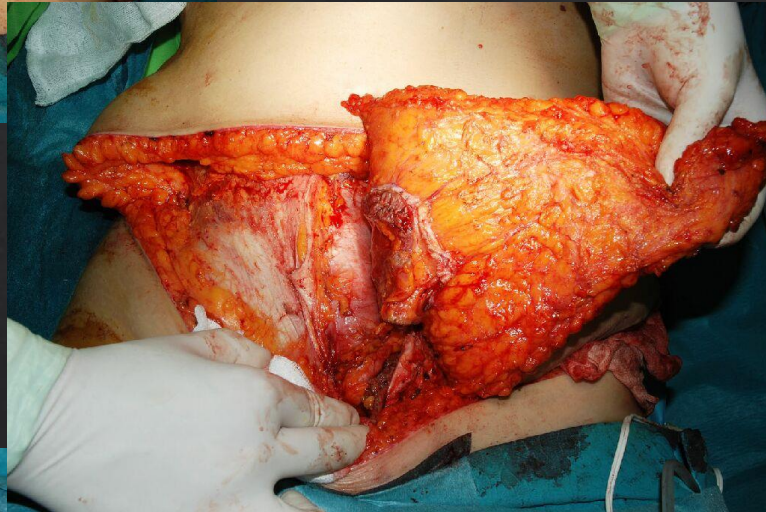


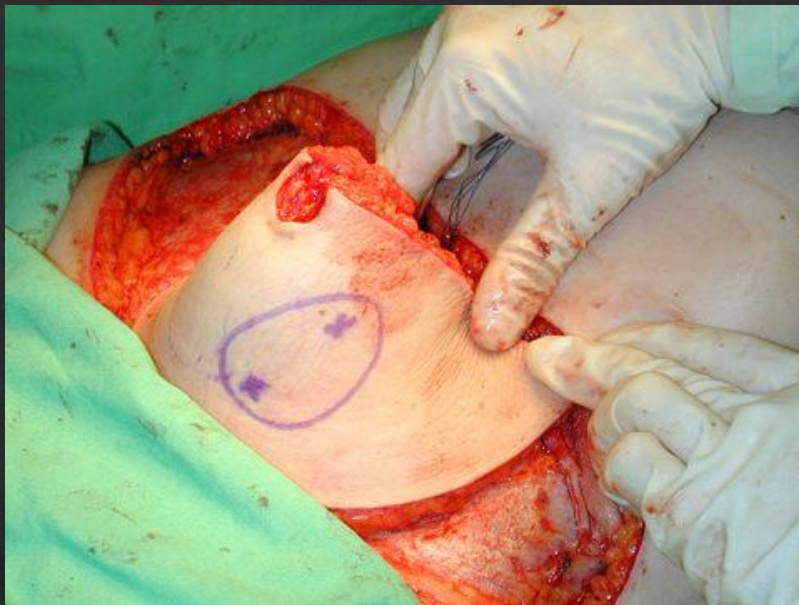
# Free DIEP flap





# DIEP





## Expander, Implantatum



# Rehabilitation

## Goal:

- familiar and economical rehabilitation
- quality of life improvement



## Type:

### • Somatic

- Pain
- Operations
- Stoma
- Voice
- Mobilitate
- Side effect of irradiation
- Side effect of chemo



### • Spiritual

- Group-therapy
- Individual therapy



# Palliation

Years and decades

**Goal:** Improving or keeping the QoL

- **Complication, pain**
- **Physical condition**
- **Spiritual condition**
- **Social condition**

- **Patient**
- **Family**



## Hospice

- **Terminal care**



# Thank You for Attention!

