



Role of Imaging in Oncology

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Basic information in Oncology

- **Tumor staging is one of the most important prognostic factors, it determines therapy** (operability, radio-, chemotherapy planning)
- **Imaging has great importance in cancer management**
 - DETECTION and EVALUATION of tumor
 - Evaluation of POST-THERAPEUTIC changes
 - Detection of the ADVERSE EFFECTs of therapy
 - FOLLOW-UP; early detection of RECURRENT TUMOR
- **Precise evaluation** is only possible with strict technical criteria, **standard protocols** and **correct image interpretation** – **our responsibility is high**

Role of imaging at the modern oncological care

Participating in the oncotherapeutic algorythm:

- TUMOR sreening
- TUMOR detecting
- Staging – T, N, M
- Therapy monitoring
- Post-therapy staging
 - Differentiation between residual tumor and scar
- Follow up
- Recurrent tumor detection - staging

–Technical expectations:

- Early diagnosis
- Precise evaluation
- Tumor extension
- Tumor size, volumen definition
- Tissue specific data collection

Imaging plays an important role also in planning radiotherapy

Imaging modalities

- **Anatomic imaging modalities** *(static/dynamic information)*
 - Conventional X-ray – mammography (digital)
 - Angiography – Digital Subtraction Angiography (DSA) – Th.
 - US
 - **CT – MD-CT** (≥ 16 detector rows in oncology)
 - **MRI** – (high magnetic field strength, 1.5T-3T)
- **Functional, molecular, metabolic imaging modalities**
 - Radioisotope diagnostic methods (bone, thyroid gl.)
 - SPECT-CT; PET/CT
 - DW-MRI, DCE-MRI, MRSI, tissue specific CA-MRI, perfusion CT, CE-US

Functional-, molecular-, metabolic imaging *imaging BIOMARKERS in oncology*

NEW measurements, qualitative, semiquantitative, quantitative
(partly in the routine examinations / partly in clinical research)

Molecular- / functional data

DW-MRI *based on: water diffusion restriction because of TU cell density, -integrity, with qualitative-, and quantitative (ADC measurement) information*

DynCE-MRI *based on: vascularisation, vascular permeability, with qualitative, semiquantitative (time-enhancement curve) information (may be also quantitative)*

Tissue specific CA *(hepatocyt-, RES specific)*

MRSI *based on: biochemical status of molecular products*

CE-US *based on: tumor neo-vascularisation*

SPECT/CT, PET/CT *(using isotope tracers, based on: different metabolic processes)*

CXR

The role of conventional radiography in the evaluation of tumor cases is limited

Today: Digital

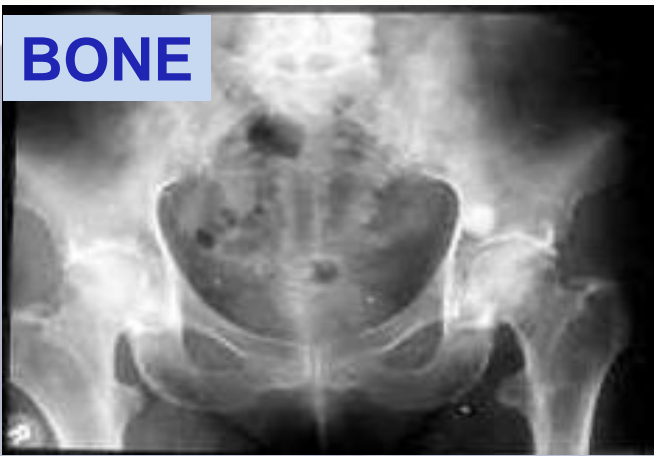
Advantages: Easy access, cheap

- Bone
- Lung
- Breast
- Abdomen
- Gastro-intestinal tract

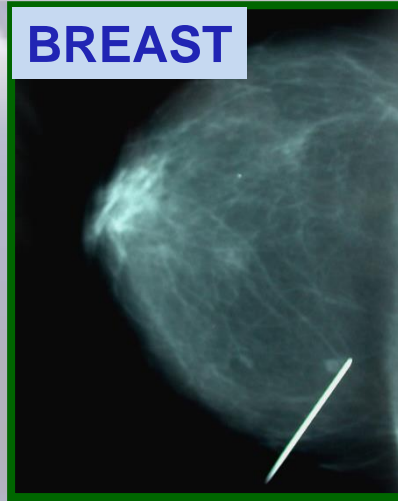
Tomosynthesis: *renewed, digital tomography for the lung and breast*

Main QUESTION: is the information enough??

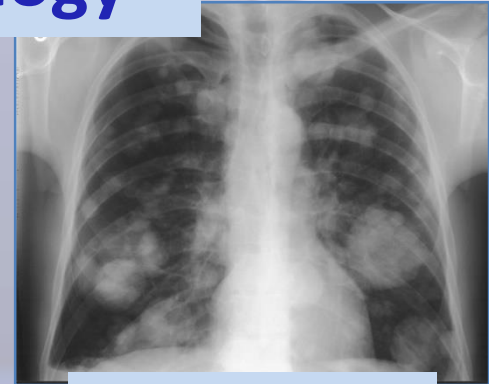
BONE



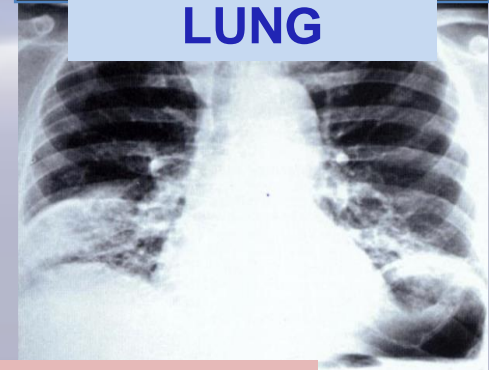
BREAST



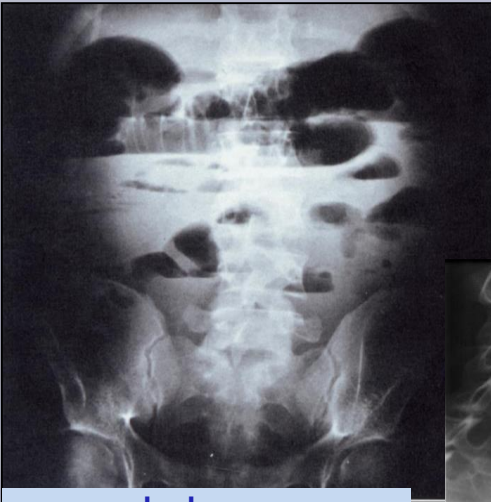
**CXR in
oncology**



LUNG



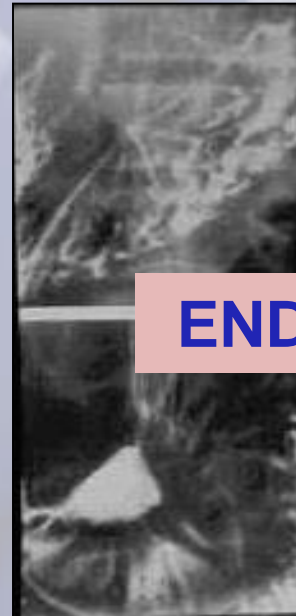
*Question:
information
will be enough ??*



abdomen
Ileus? perforation?

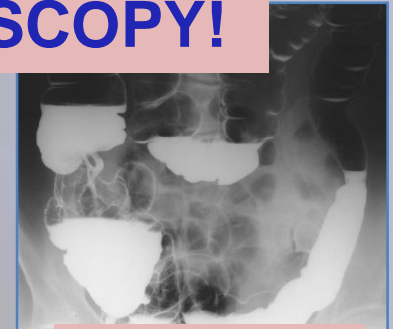


esophagus
swallowing function



stomach

ENDOSCOPY!



colon

Ultrasonography

excellent for the soft tissue

Advantages:

- Easy access, cheap
- Non invasive, non ionising
- Tolerable
- Real-time information

Clinical applications

- Transcutan – abdominal, pelvic, neck, breast, extremities
- Vessels
- Endocavital, - rectal, - oesophageal, - endoscopic US
- Intraoperative US
- US guided biopsy/drainage

Disadvantages :

- Lack of complex information
- Difficulties in the evaluation of
 - Deep structures
 - Big lesions
- Subjective
- Hard to standardize

Methods

- Gray scale
- Doppler
- **CE-US**
- US-elastography

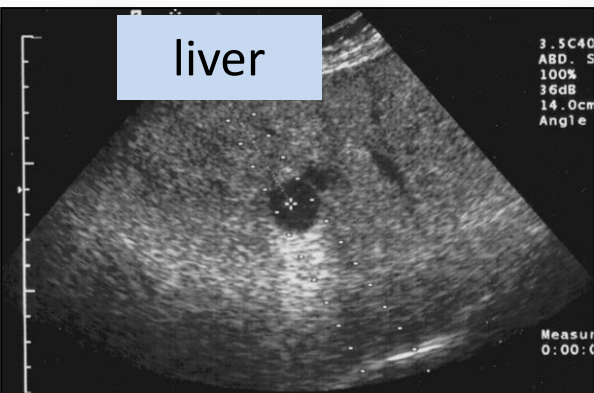
CE-US - HCC

arterial, portal, venous, parenchymal phases



**US is not the standard tool
for tumor evaluation**

liver



US
excellent soft tissue
resolution

BUT
lack of complex
information

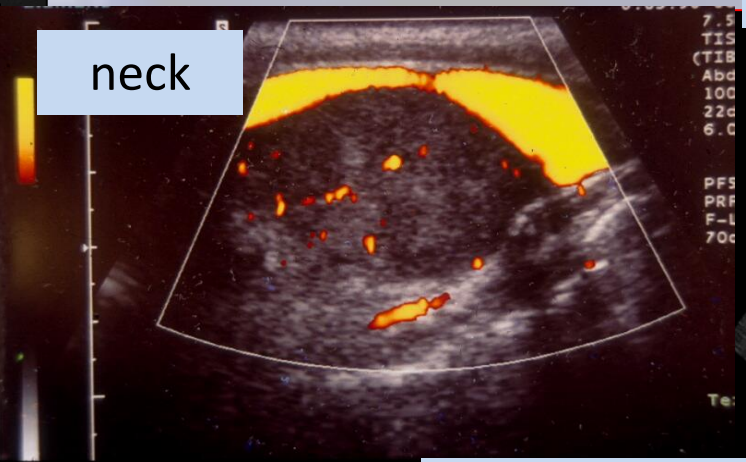


Endorectal US-in rectal ca

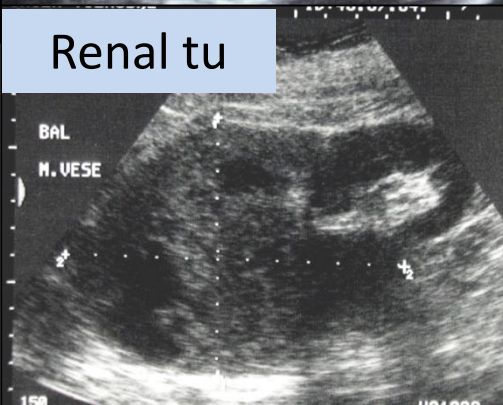
Bile duct



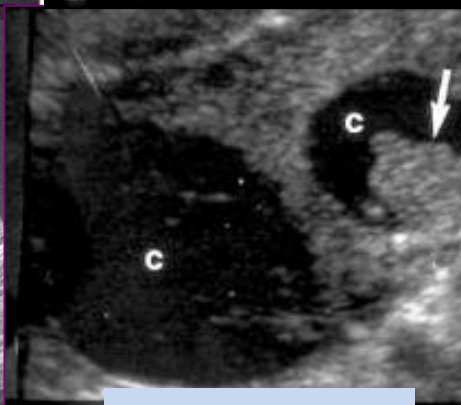
neck



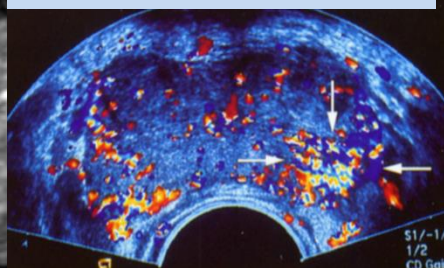
Renal tu



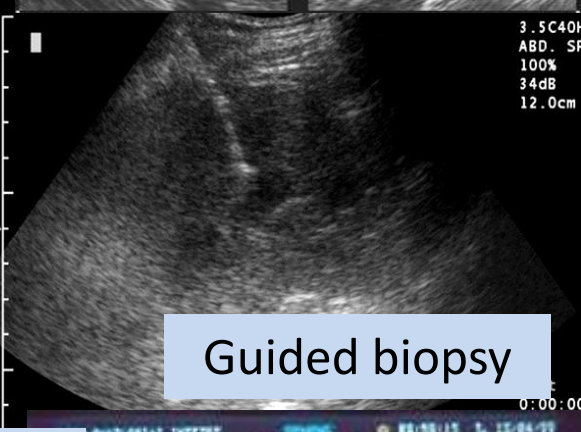
Endorectal US-
in prostate ca



Ovarian ca



Guided biopsy



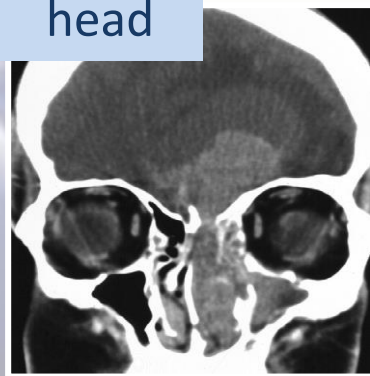
CD-US

Advantages of MD-CT

- Quick, tolerable, informative, standardized
- Whole body information
- Optimal vascular phase settings
- Dynamic / Perfusion information
- Excellent multiplanar reconstructions and 3D images
 - CTA & virtual endoscopy
 - We can see better, more and detect smaller changes
 - More structural details

**Disadvantage:
ionising
radiation**

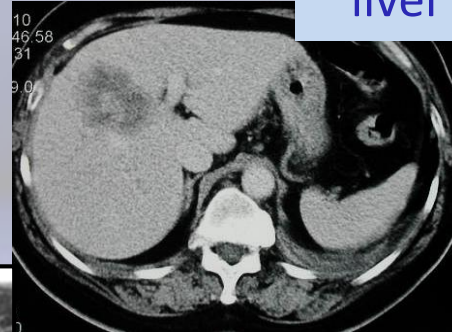
head



CT

Quick, informative, standard method

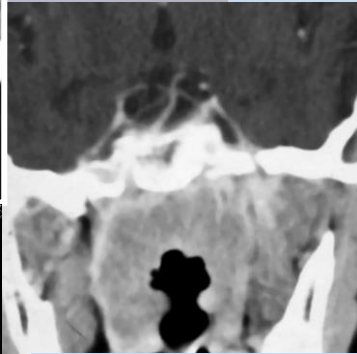
liver



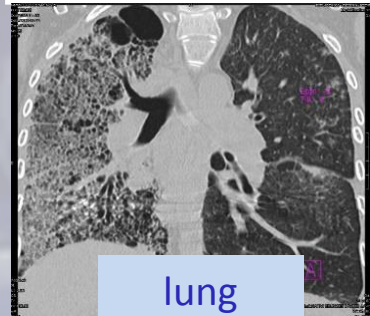
Guided biopsy



Skull base, neck



lung



lung

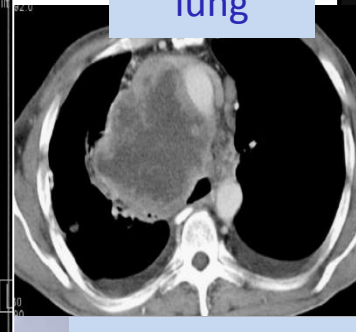


pelvis



Guided drainage

mediastinum



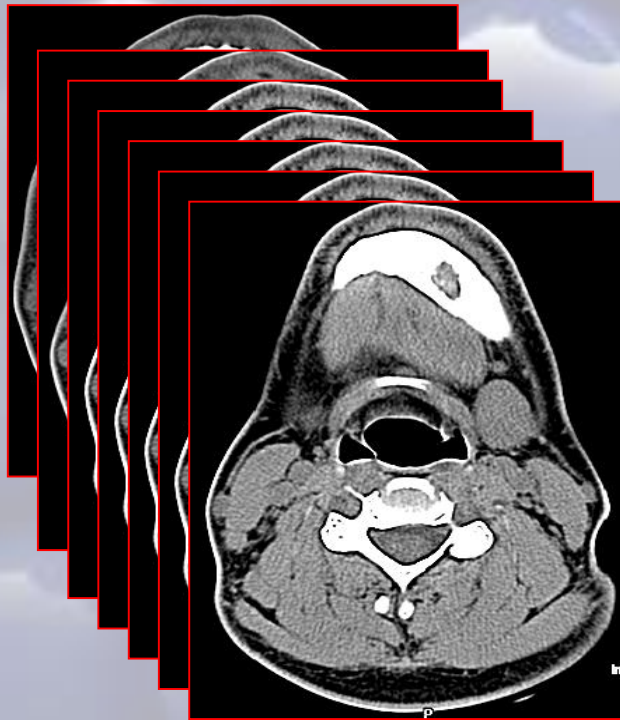
Bone, spine



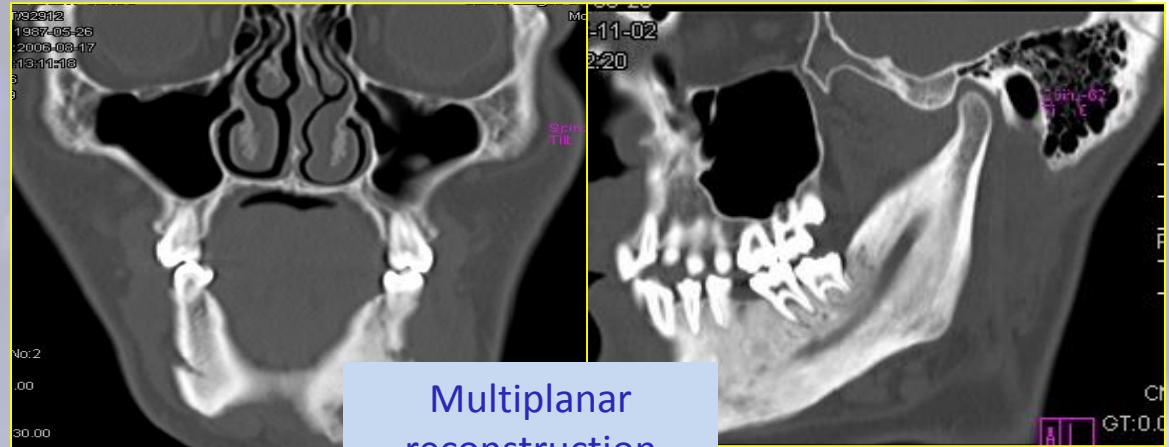
Whole body



MD-CT -Volumetric measurement – Multiplanar-, 3D information



Volumetric
measurement



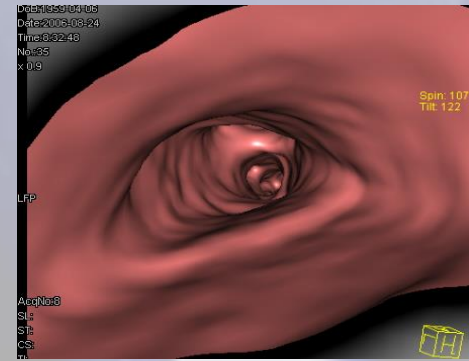
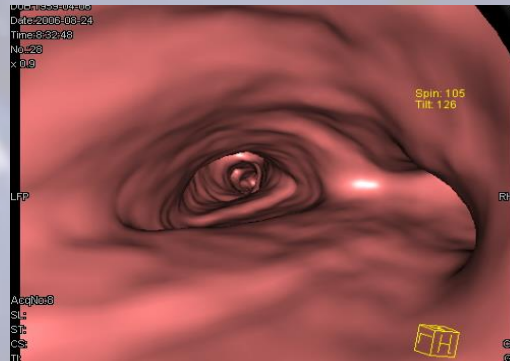
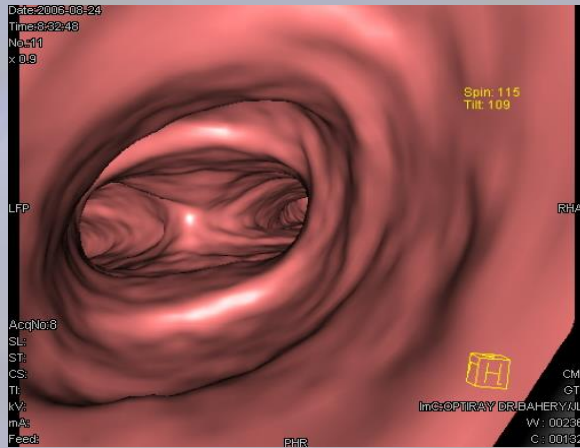
Multiplanar
reconstruction



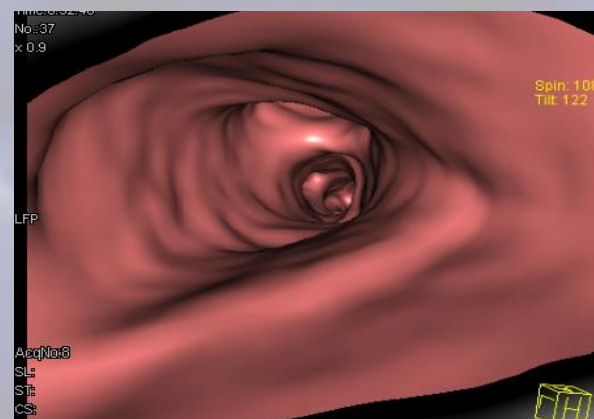
3D information

MDCT

Virtual endoscopy



based on volumetric data collection



CT- Angiography

Magnetic Resonance Imaging- MRI ($\geq 1.5\text{T}$)

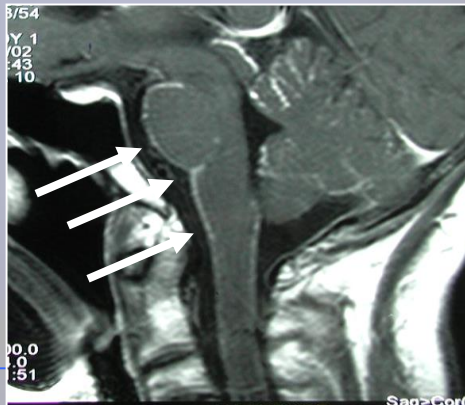
excellent multiparametric modality

with high spatial & high contrast resolution

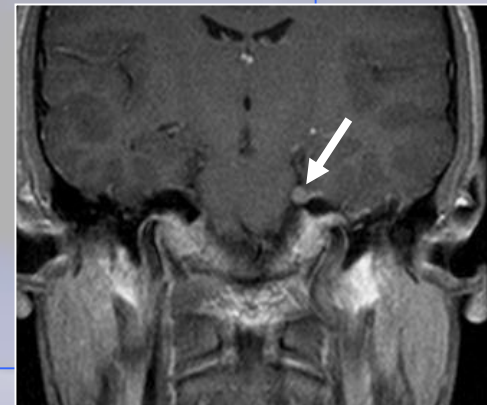
- **Best soft tissue evaluation** of intracranial-, perineural spread, spine, head and neck, pelvis, upper abdomen, breast, extremities
- **Tissue specific information:** fat, melanin, blood, etc. Extracellular-, hepatocyt-, RES-specific contrast agents
- **Functional information:** diffusion-weighted MRI (DW-MRI), dynamic contrast enhanced MRI (DCE-MRI), MR-spectroscopy (MRSI)
- **Flow sensitivity**
 - MR angiography

**without
ionising
radiation**

ATTENTION!
Methal within the body!



Lepto-meningeal TU spread



Perineural (N.V.) TU spread

MRI „gold standard“

Problem solving method!

BETTER than CT at:

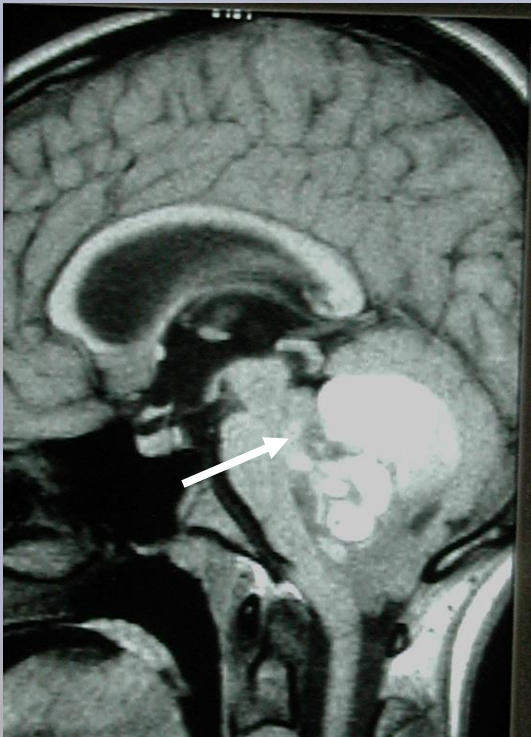
- Brain tu: assessment of Tu volume/localisation/function
- H&N staging: intracranial / perineural tu spread; Igl evaluation
- **Abdomen** (*fast sequences*): **LIVER**; *pancreas, kidney, adrenal gland, GI*
- **Pelvis**
 - Prostate ca / Gynecological tu / Rectal ca
- Bone (marrow) – metastasis
- Soft tissue tumors

MRI ACC: >80-90%

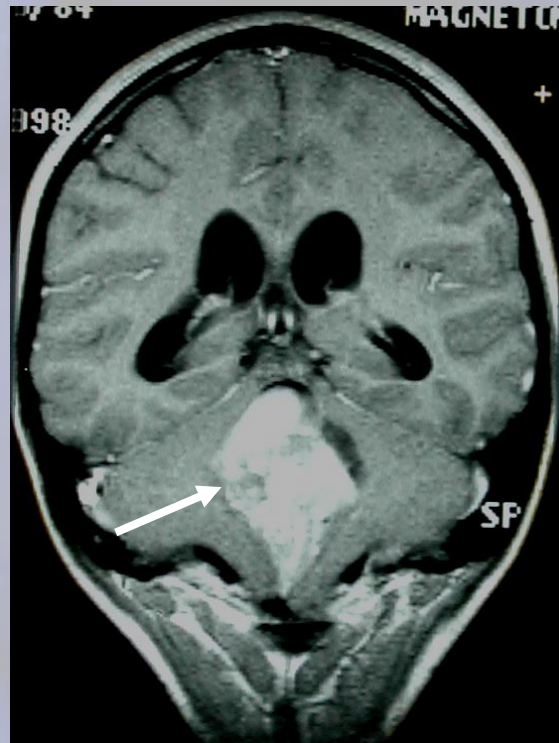
MEDULLOBLASTOMA in the IV. ventricle

MRI - CE-T1-w images

Best multiplanar evaluation of intracranial tumors



Sagittal

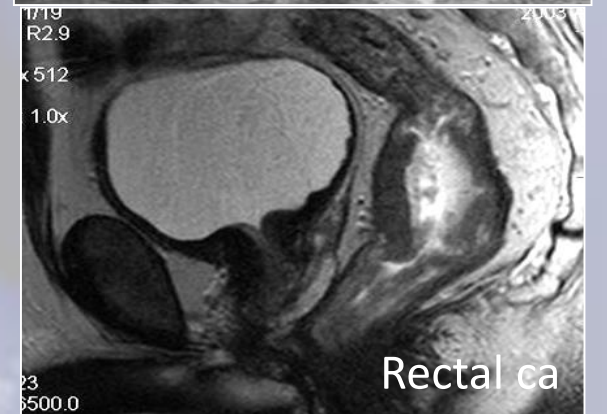
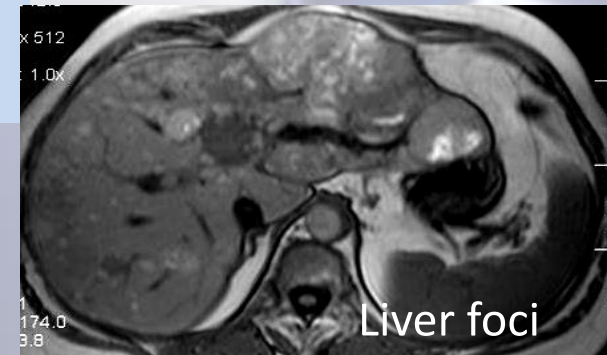
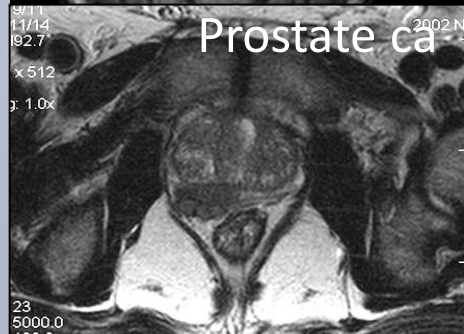
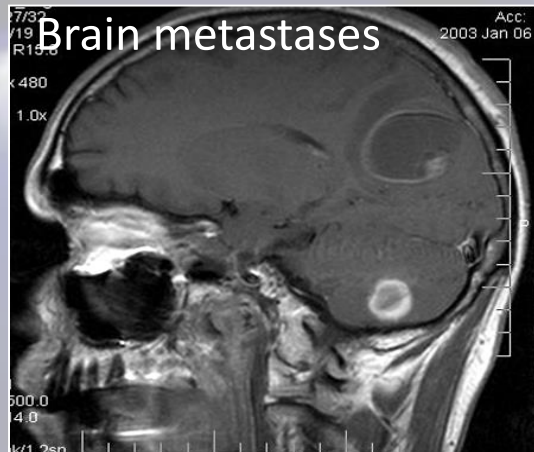


Coronal



Axial

Today: MRI - Basic method



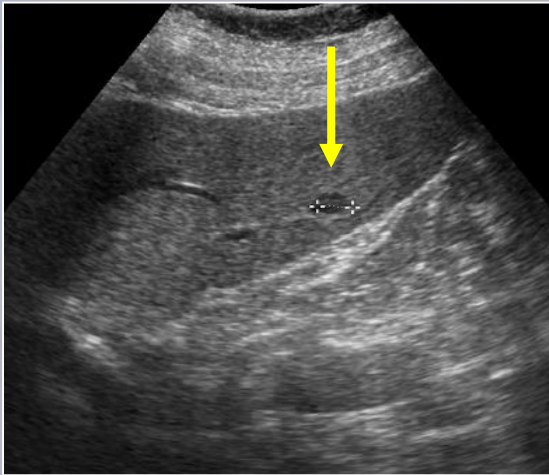
Tissue specific information

Two malignant primary tumors

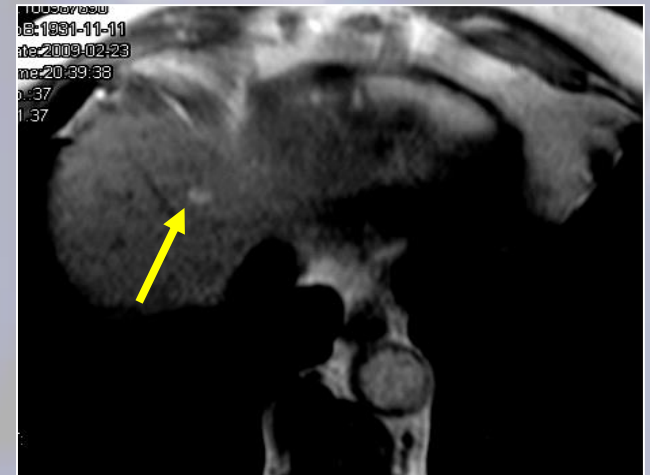
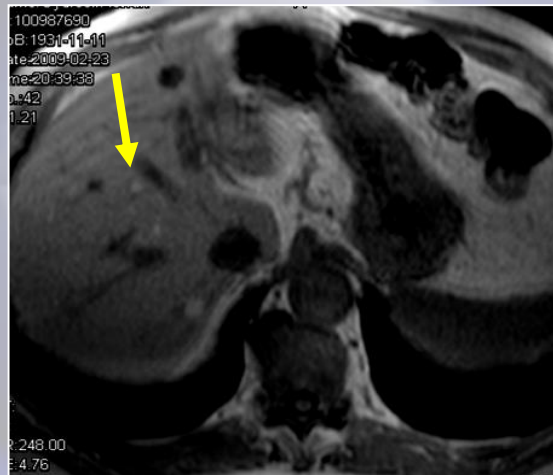
Colon ca / ocular malignant melanoma

MRI: specific for MM metastasis

*High signal intensity T1-w foci in the liver -
because of melanin content*



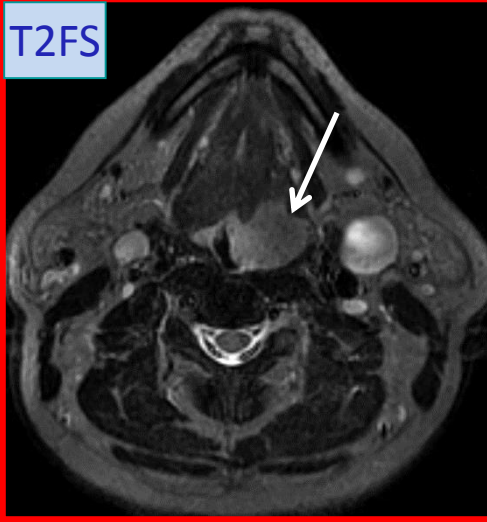
US – unspecific density
It might be metastasis



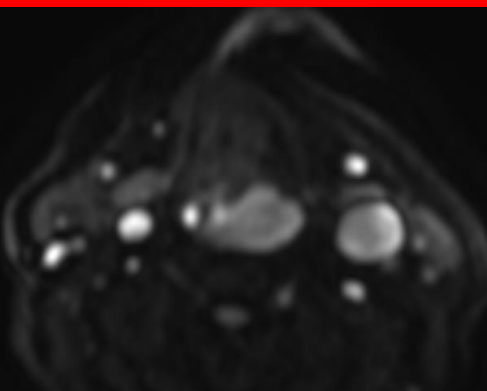
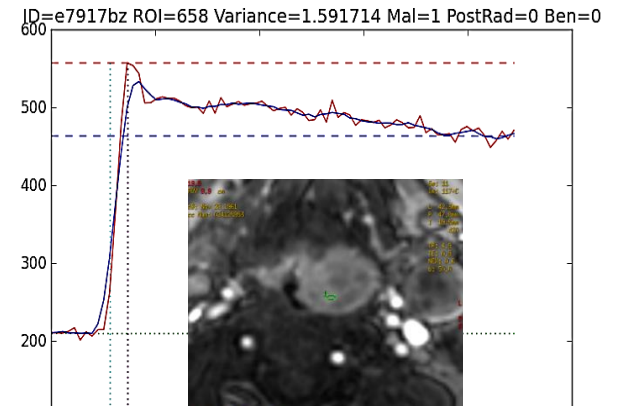
DG: MM mets

MP-MRI – anatomic and functional measurements in mesopharynx CA (native T1-,T2-w, CE-T1FS, DW-, DCE-MR)

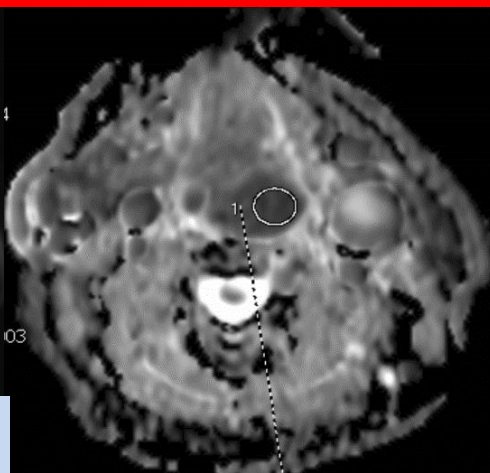
T2FS



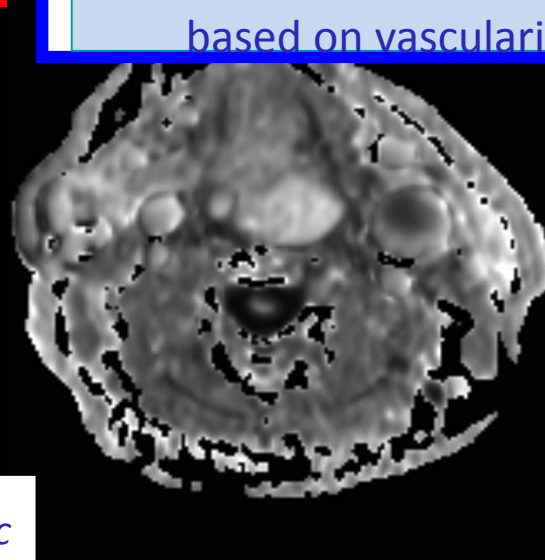
CET1FS



DW-MRI: based on TU
cell density
b-value: 1000 s/mm²



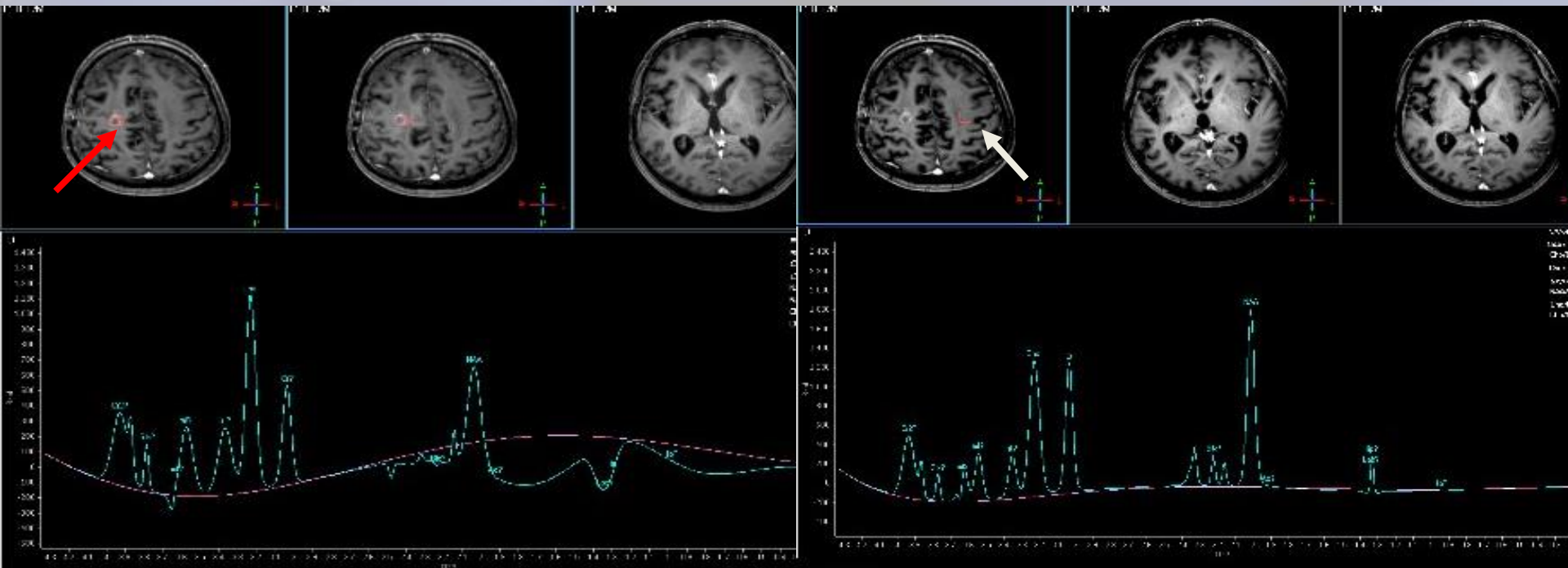
ADC: $0.743 \times 10^{-3} \text{ mm}^2/\text{sec}$



MR spectroscopy (MRSI) – (biochemical analysis of *molecular products*) *Recurrent brain tu- could be detected earlier*

Tumor side (R)

Normal side (L)



**Cholin
peak**

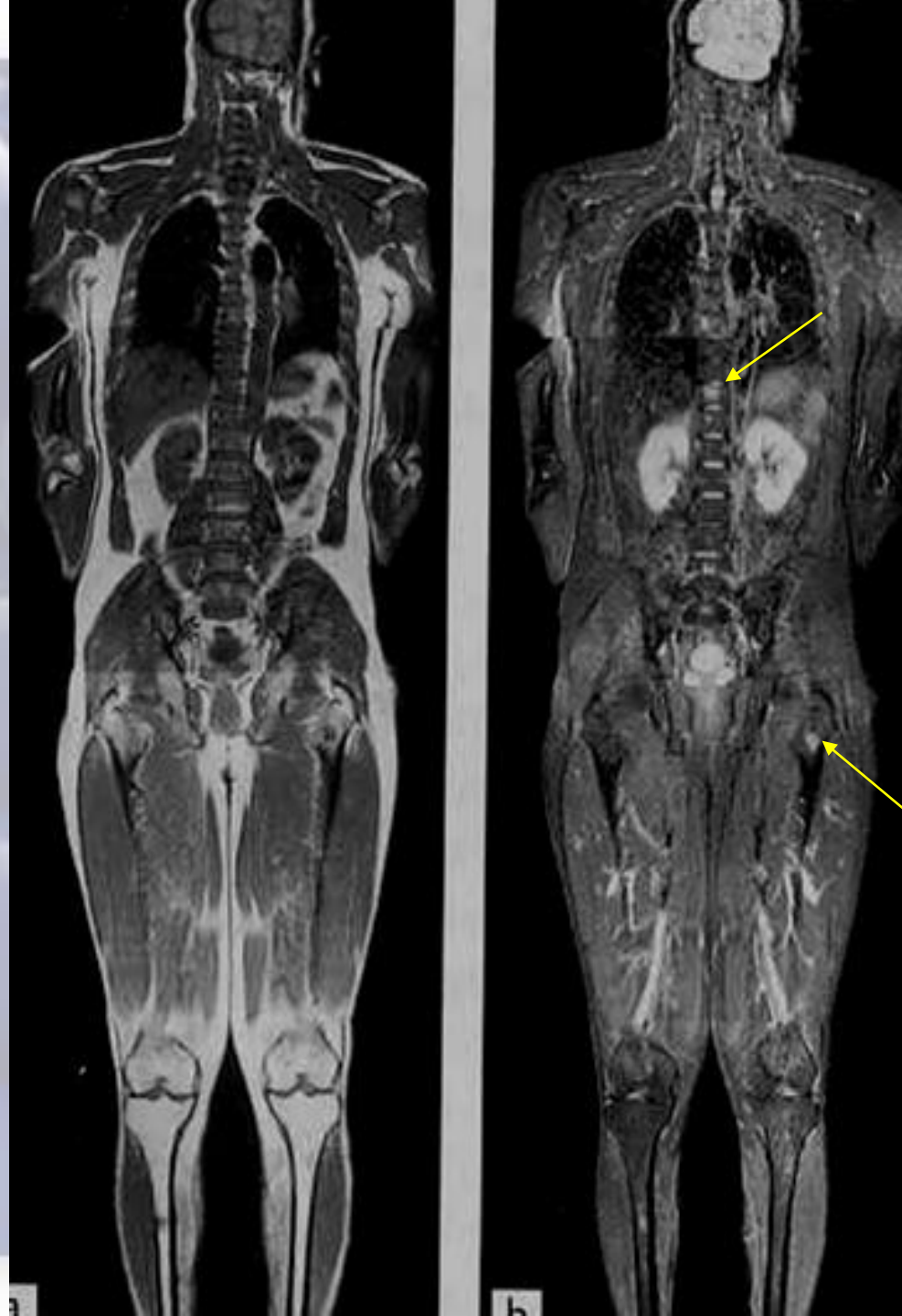


NAA ↓
N-acetylaspartate

Cholin ↓

NAA ↑





Whole body MRI

Sensitive and specific for
bone marrow metastasis

Right: T1-w sequence

Left: STIR sequence

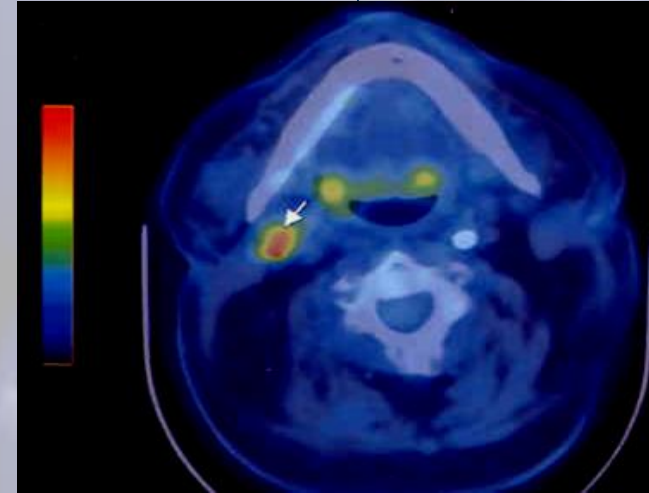
PET/CT

- **PET/CT – *better imaging of molecular processes***
- Whole body information
- **PET: sensitive for metabolic activity – Tracer: FDG**
(F18FluoroDeoxyGlucose) – glucose alternative
- **CT: anatomic background and structural analysis**
- **Together: improved sensitivity/specificity/accuracy**

PET/CT

general indications

- Clinically suspected distant metastases
- Therapy monitoring
- Assessment of residual / recurrent tumor
- Re-staging in case of recurrent disease
- Unknown primary (cervical metastatic lymph node – detected primary tumor in 30-40%)



False NEGATIVE & POSITIVE cases!
Ionising radiation!
High costs!

FDG-PET/CT

Two primaries

1. Radix linguae + N met
2. Non-Hodgkin-Lymphoma in the abdomen

01-22
15

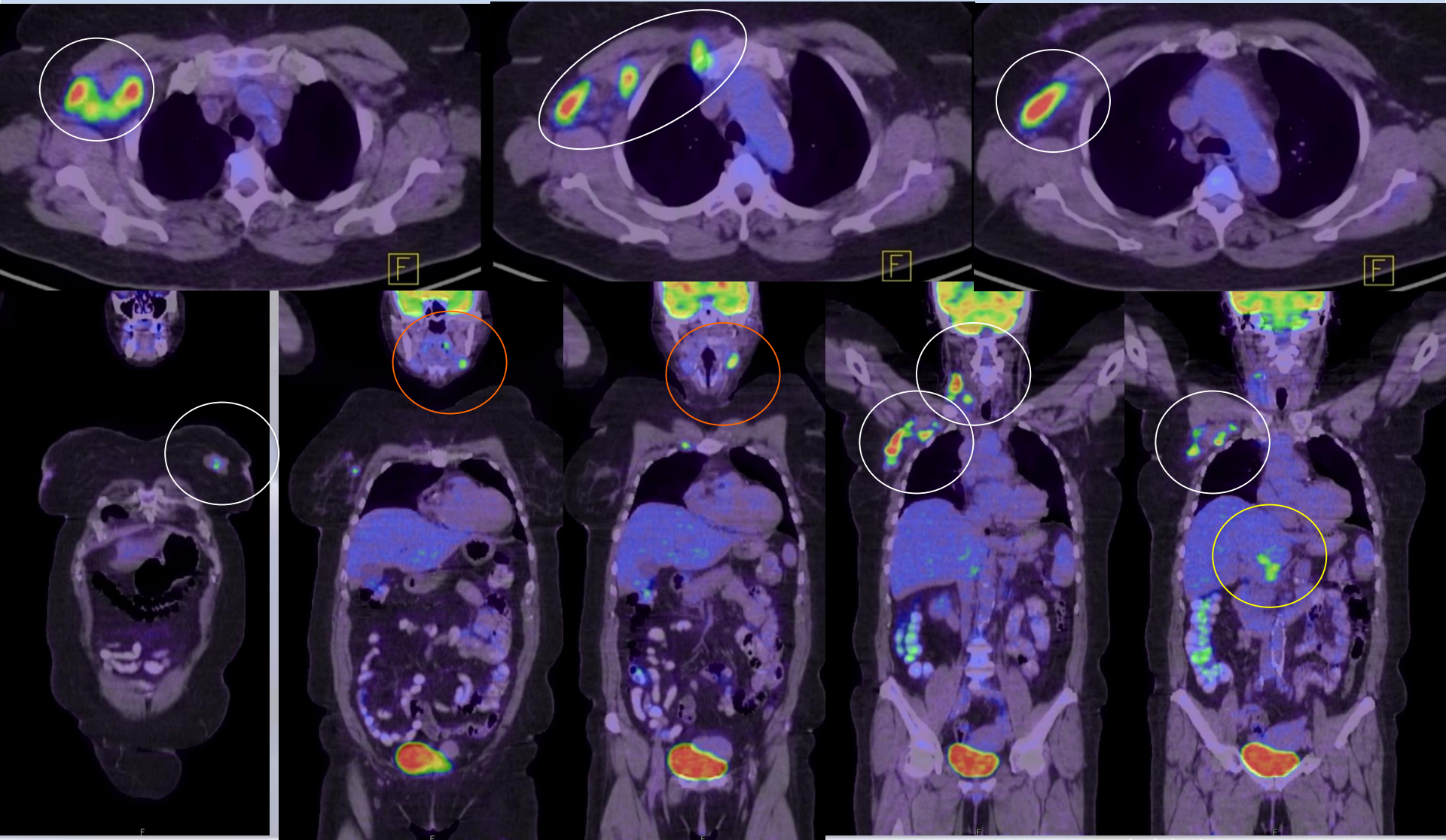
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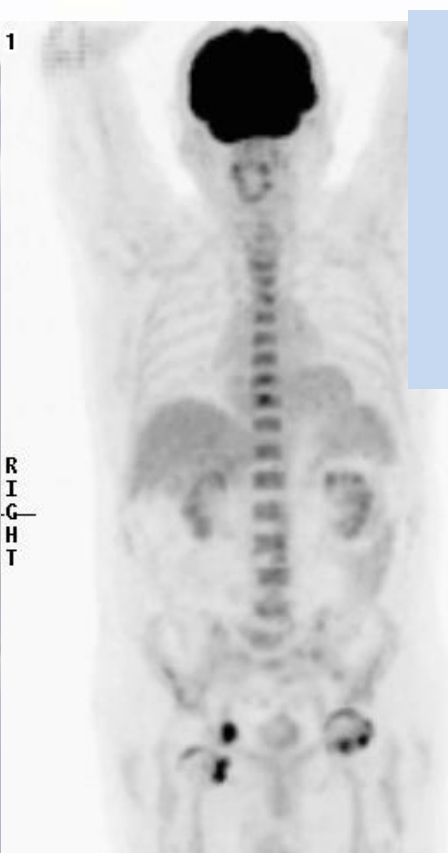
22

FDG-PET/CT – whole body information

three primary tumors

(left mesopharynx-, right breast-, cholangio ca)





**RN-
osteoscan**
Tc-99m diphosphonate,
Very sensitive, but less
specific

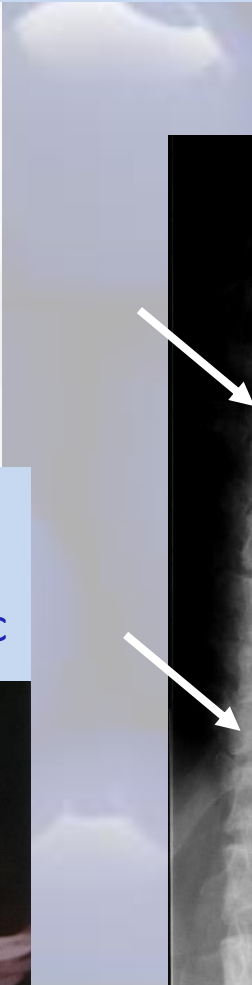
based on: osteoblast activity

BONE
Spine
metastasis



MR
Sensitive and
specific

CXR
Specific but NOT
sensitive



CT
Medium specific and
sensitive



PET/CT
sensitive and specific



Interventional radiology in oncology

Diagnostic

➤ Diagnostic angiography - DSA

vascular morphology,
neovascularisation, cancer
vessels

➤ Guided biopsy

(Fluoroscopy-, Mgr-, US-,CT-,MR-)

- FNAB – fine needle aspiration biopsy for cytology
- core biopsy for histology

Therapeutic

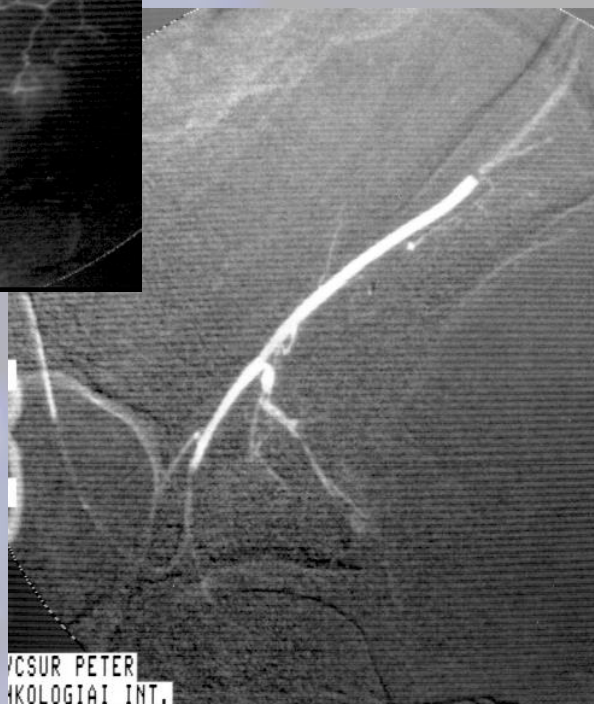
- Intravascular therapy - DSA
 - TU embolisation,
 - TU chemoperfusion
 - Dilatation, stenting
- Tumor ablation (with radiofrequency-, (RFA) Laser wave, percutan ethanol injection (PEI), focused US)
- Drainage (*abscess*)



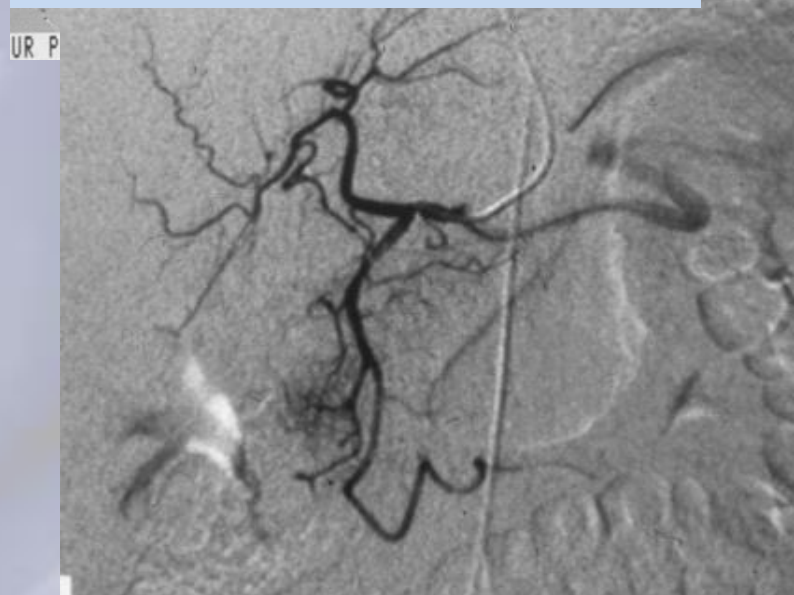
DSA -TH
Localized cancer



Chemoperfusion

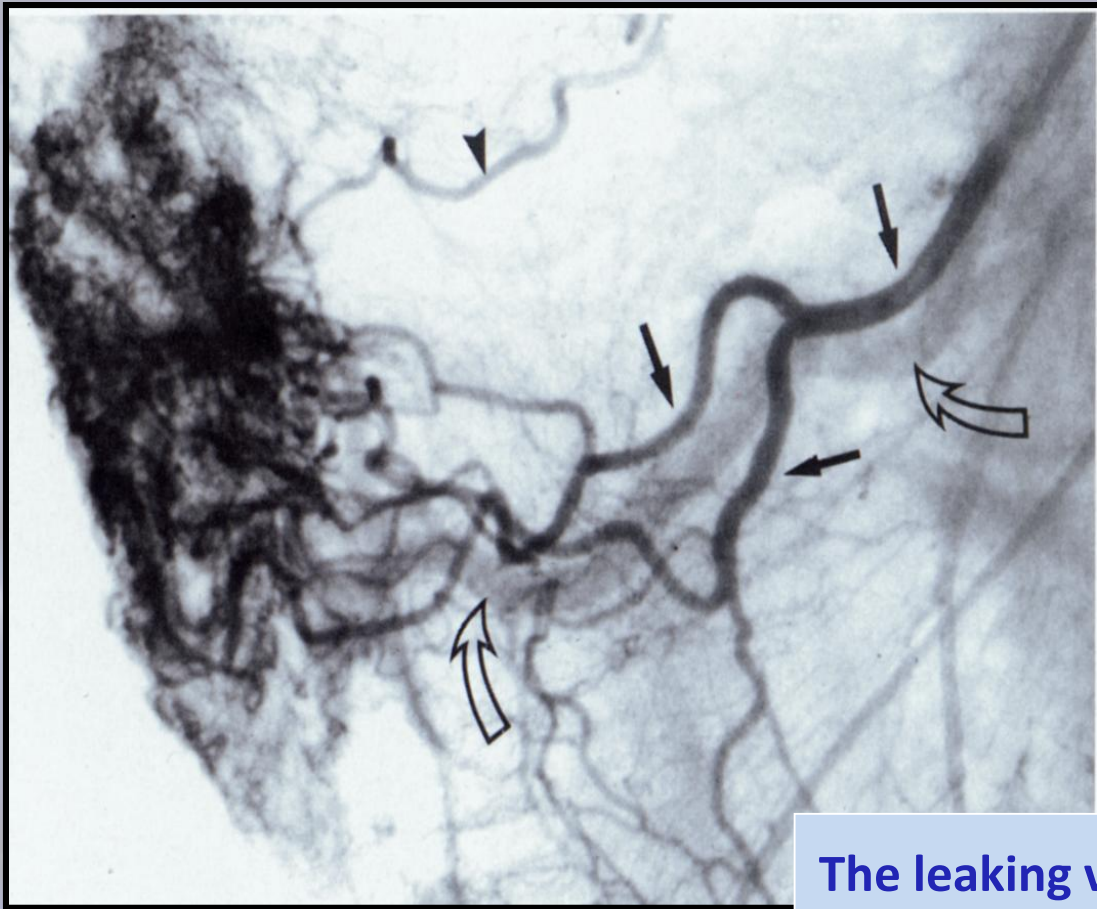


Chemoembolisation
Cancer vessels have been closed



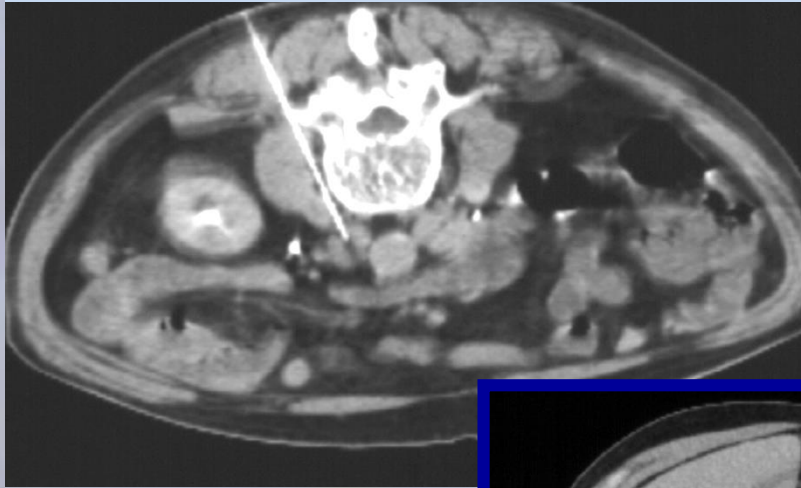
Cancer vessels were demolished

Embolisation of coecal AV malformation - because of bleeding -



The leaking vessels were
obstructed

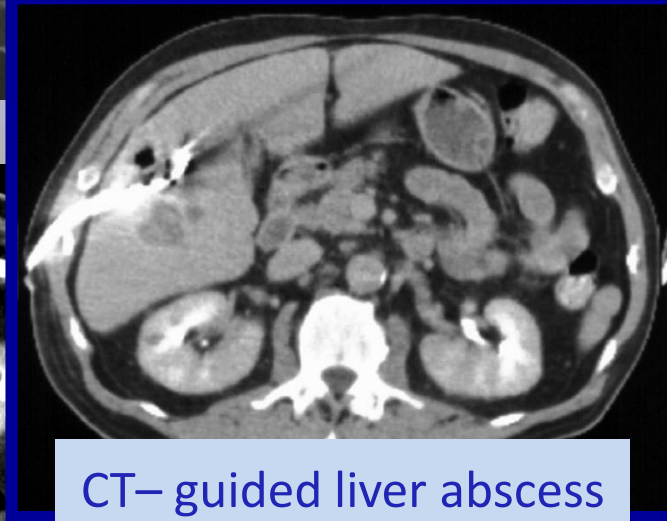
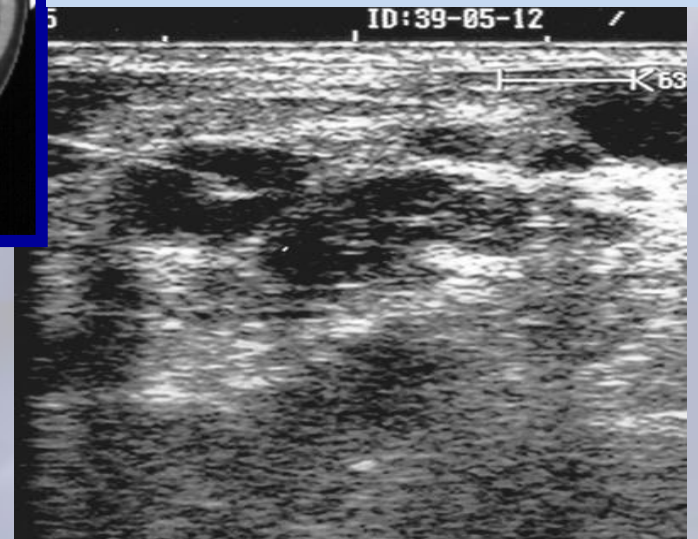
CT– guided lymph node biopsy



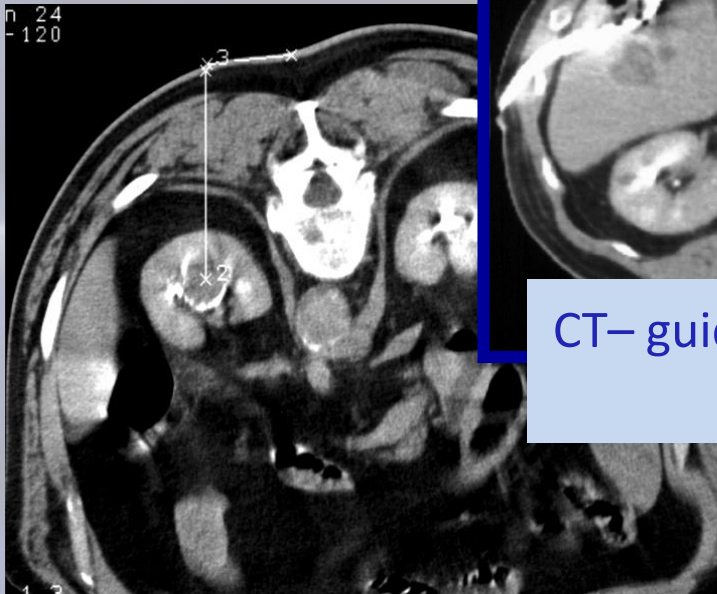
US, CT– guided biopsies,
drainage



US – guided neck node biopsy



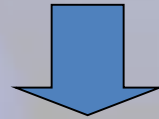
CT– guided liver abscess
drainage



CT– guided renal biopsy

Oncological diagnostic algorithm

**DETECTION
SCREENING**



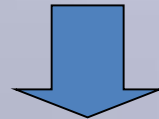
XR, US, CT, MRI
biopsy / guided

STAGING



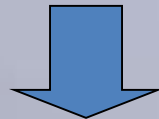
CT, MRI, RN, PET, biopsy /US/CT guided

**THERAPY
RESPONSE**



CT, MR, RN, PET, US, XR

FOLLOW UP



US, CT, MRI, RN, XR.

**RECURRENT TU
RESTAGING**

CT, US, MRI, RN, PET

Requirements of SCREENING

Rational chance for:

- Early diagnosis in preclinical stages
- To find high risk asymptomatic individuals
- To achieve lower mortality rate

*For example: 90% of all **breast cancer** can be cured with early diagnosis and appropriate therapy!*

Mammography

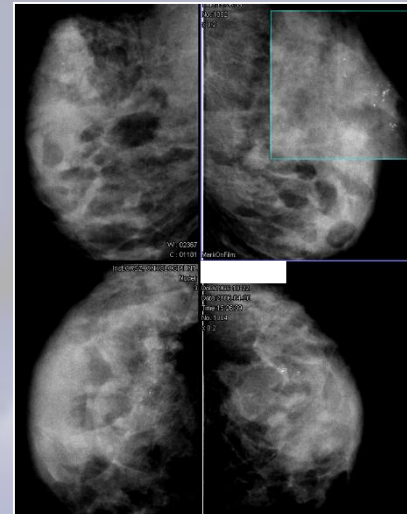
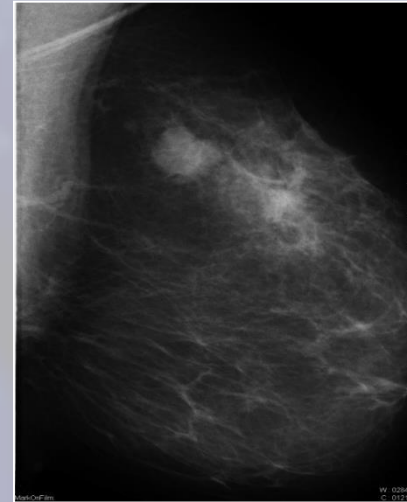
- basic screening method for breast cancer -

SENSITIVITY

- Average in literature: 80-85%
- In adipose breast: 99%

As breast density increases
sensitivity decreases!

For dens breast: additional US, MRI



BREAST CANCER

Stage determining imaging methods

- ❖ **Mammography - Analog / Digital**
 - ❖ Tomosynthesis
 - ❖ CAD (Computer Assisted Diagnosis)
- ❖ **US**
- ❖ **Guided biopsy:** FNAB / core-, vacuum assisted biopsy
by US / mammography (stereotactic biopsy)
- ❖ **Multiparametric MRI** (Anatomical and functional assessment)
- ❖ **CT / PET-CT – for staging (metastases?)**
- ❖ **Localization before op.:**
 - a) Radioguided localisation(ROLL) for occult lesion, SLNB
 - b) Hookwire-guided localization for non-palpable breast lesions
- ❖ **Specimen** mammography /US

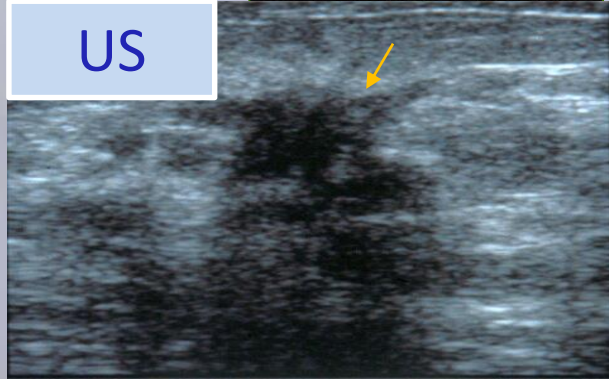
BREAST CANCER

Multimodality - Multidisciplinary

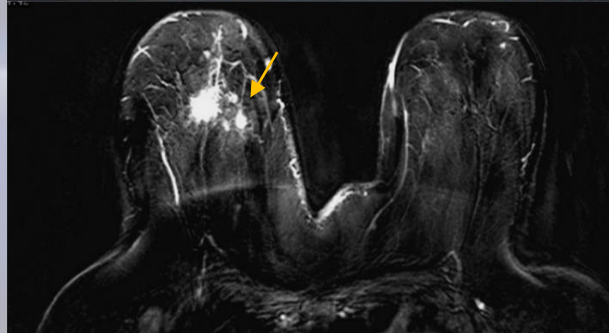
X-ray-mgr



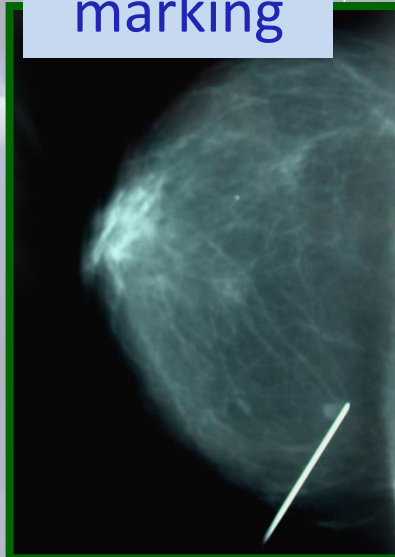
US



Breast MRI



marking



Mammography + US + biopsy

Sv 85%, Sp 92-95%

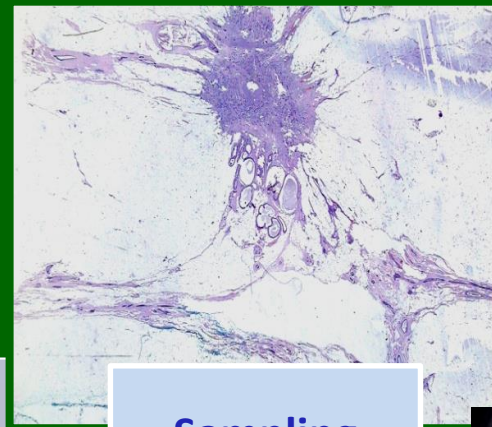
Breast MRI:

Sv 95%, Sp 86%

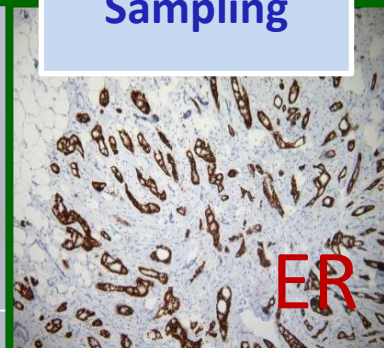
Sentinel N

Lymphoscintigraphy +
+ (Blue dye)
+ Histology
(Accuracy >90%)

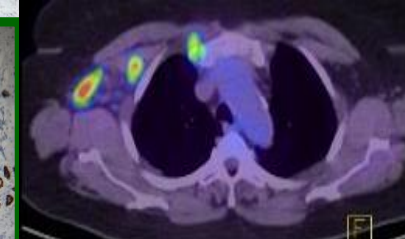
Sampling



ER



CT, PET/CT
staging



T/N: mammography / US / MRI /+sentinel N

LUNG CANCER

- **Leading cancer death**
 - 1.3 million deaths / year worldwide
 - >60.000 deaths in 2010 in USA
 - Approximately 70% of cases are incurable at presentation, metastatic or locally advanced
 - **14% overall 5 year survival**

Theresa C. McLoud, MD
Massachusetts General Hospital, Harvard Medical School

LUNG CANCER SCREENING

- **CT - highly sensitive for small lung nodes**
CT - detects more cancers than CXR
- **CT screening for lung cancer has mortality benefit**
NSCLC: in Stage IA - T<3cm, N0, M0 - survival > 65%
- T<1cm, N0, M0 – survival > 80%
(Henschke study)
- **Low dose CT** (minus 20-25% of standard dose)
 - Follow up LDCT for grow
 - Volumetric measures - CAD (computer assisted diagnosis)
- **Risk:** currently no determined risk limit at CT screening for lung cancer
- **High risk group:** smokers >1 pack/day; >55 years old
(Dr. Lecia V. Sequist, Massachusetts General Hospital, Boston)

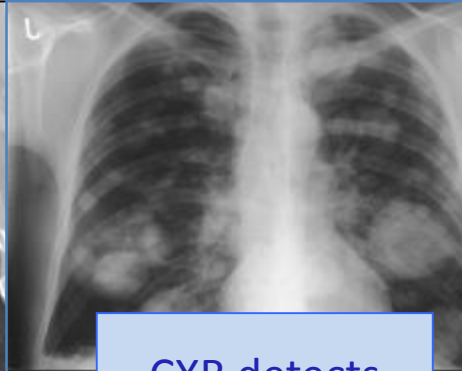
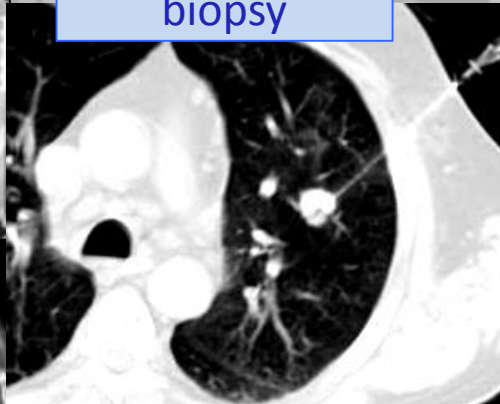
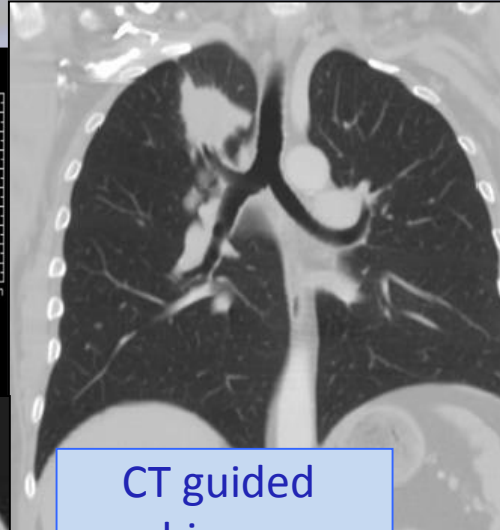
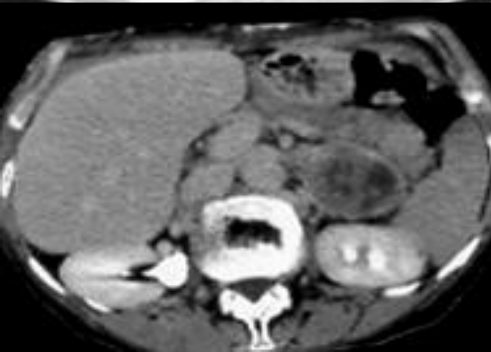
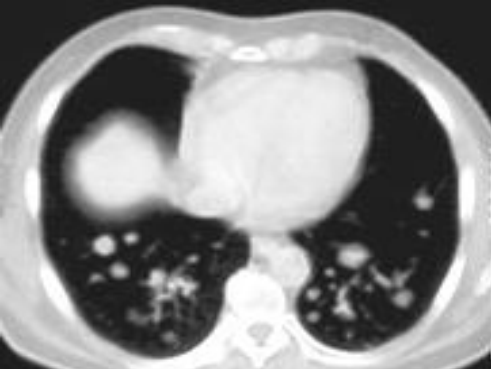
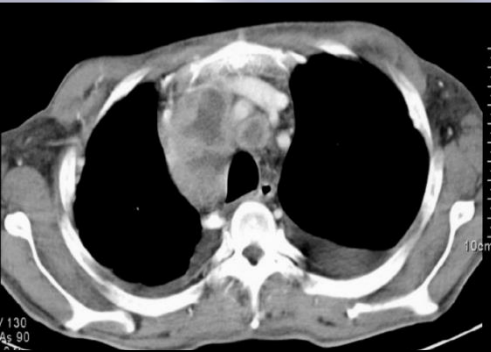
CT basic method

- Staging-
- T-Acc 90%

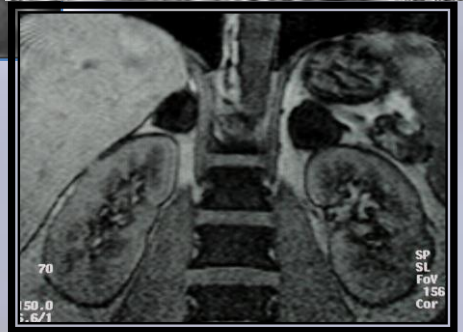
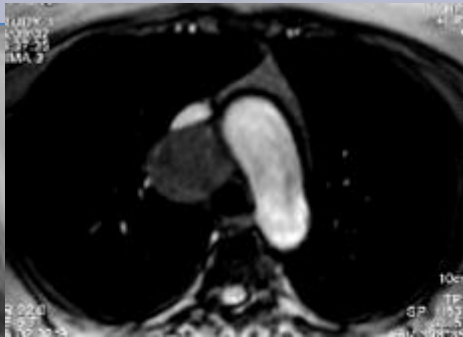
LUNG CANCER

Clinical exam.: Bronchoscopy

Complementary
MRI



CXR detects



PET/CT
staging

Lymph node &
adrenal gl. met

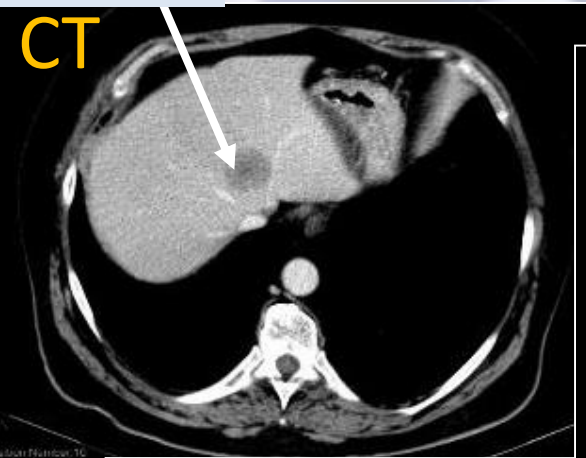


LUNG CANCER METASTASES

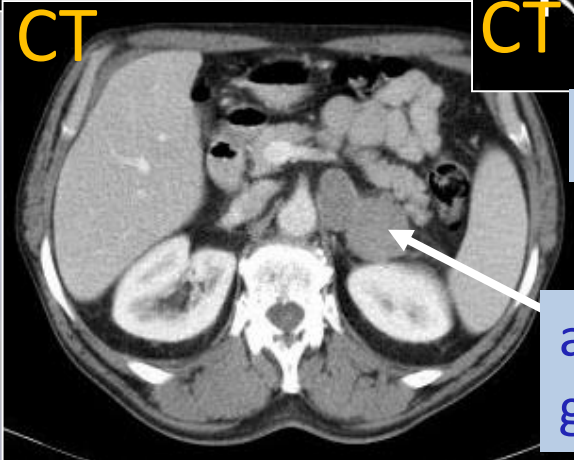
(CT / MRI / PET-CT)

liver

CT

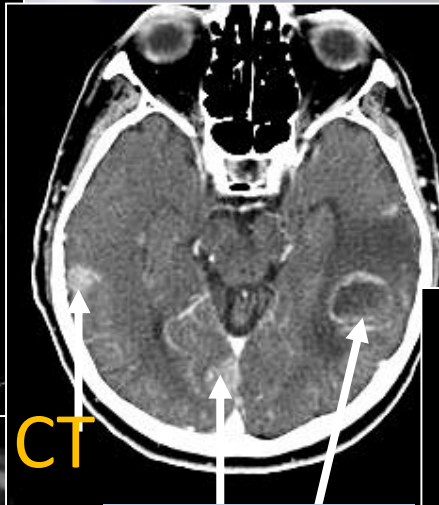


CT



adrenal
gland

CT



brain

PET/CT

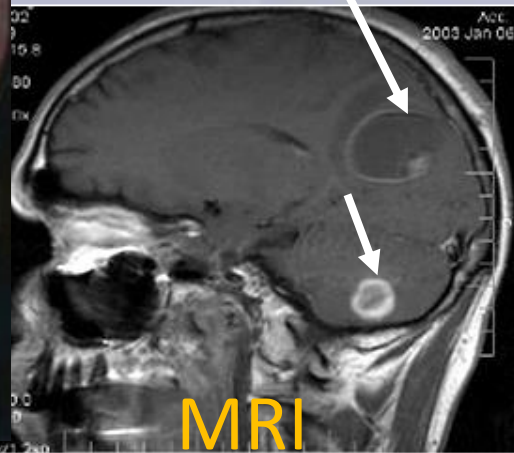


CT

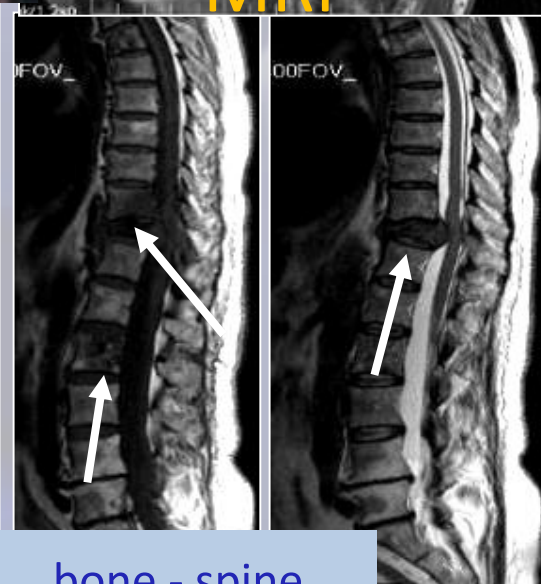


brain

MRI



bone - spine



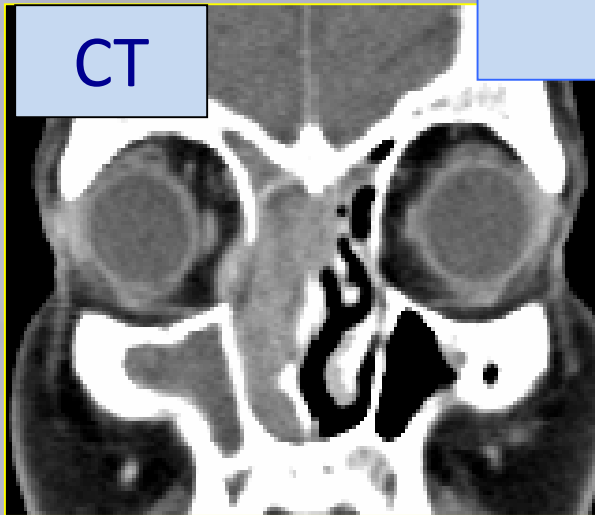
Imaging in **HEAD** and **NECK** tumors

- **US** – for analysing palpable neck masses
 - solid / cystic? , lymph node, thyroid gl., salivary gl., vessels
 - Guided biopsy
- **CT** (*complete imaging of neck from the skull base to the trachea bifurcation*) + facial bones
- **MP-MRI**: best modality to evaluate the local staging
- **PET/CT** - for whole body information – for distant TU spread, for residual /recurrent TU

Head & Neck Ca: MR/CT/US

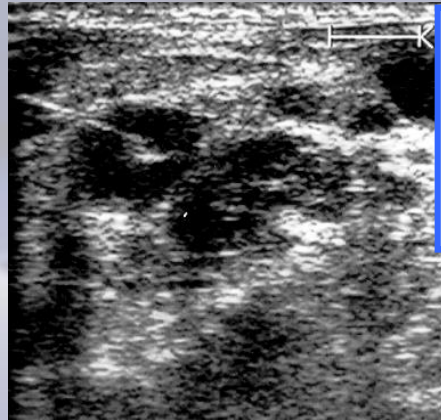
Clinical examination: endoscopy

CT

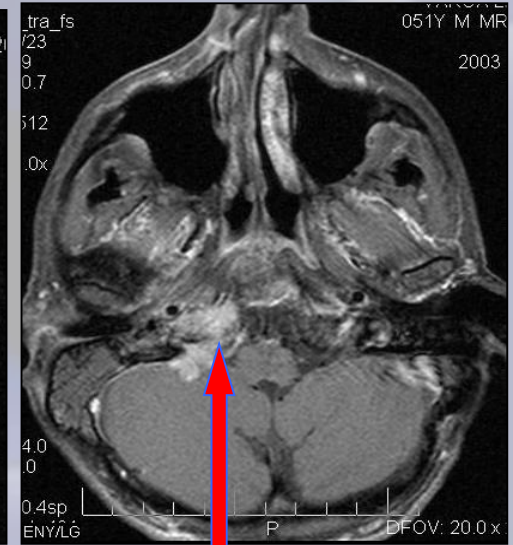
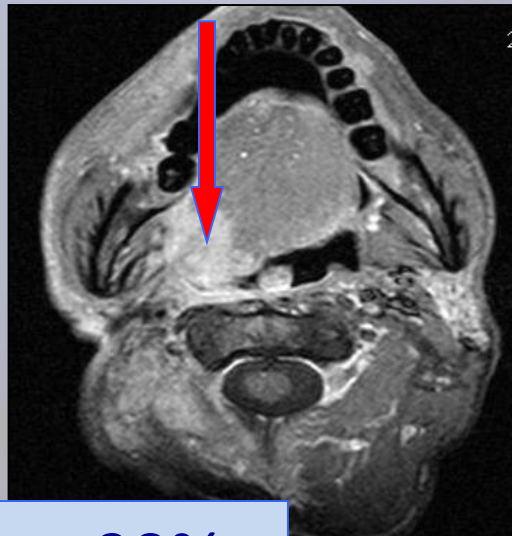
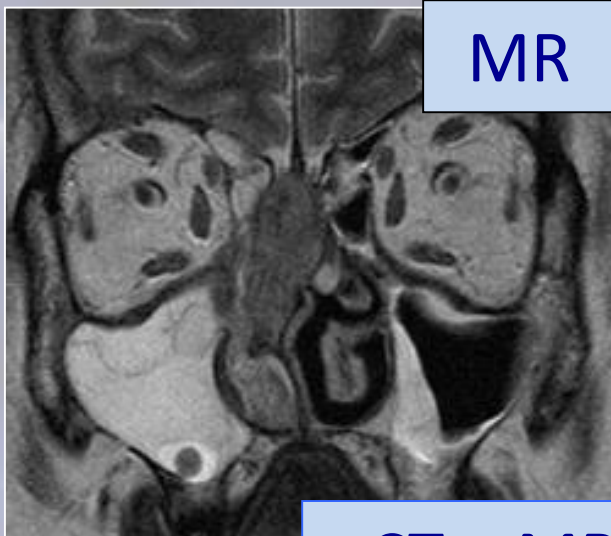


US

Guided
Asp. Cyt.
N - Acc >90%

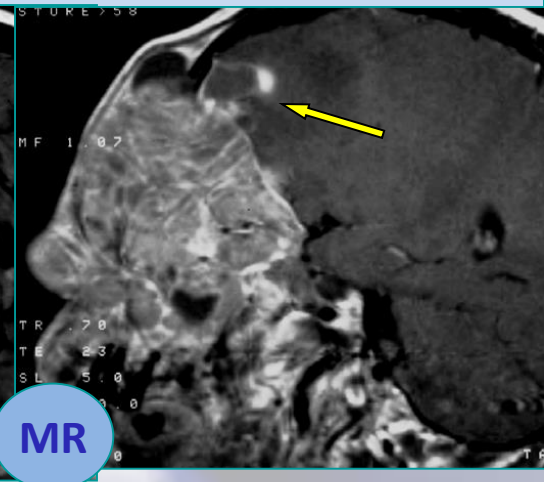
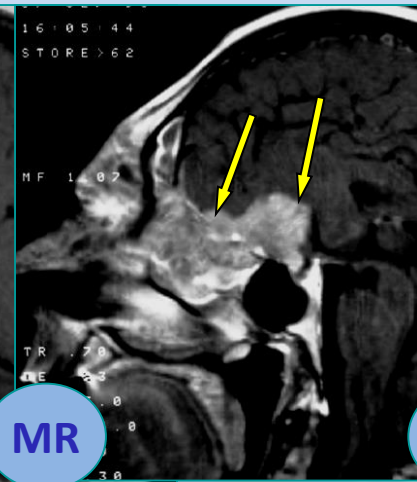
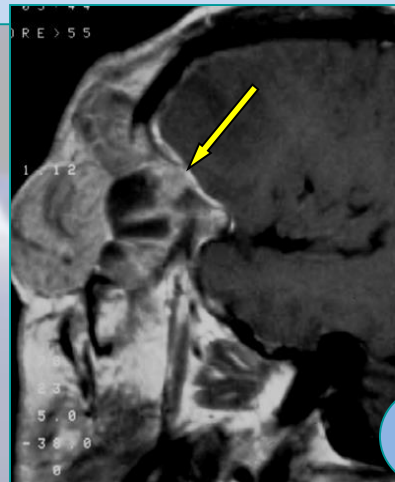
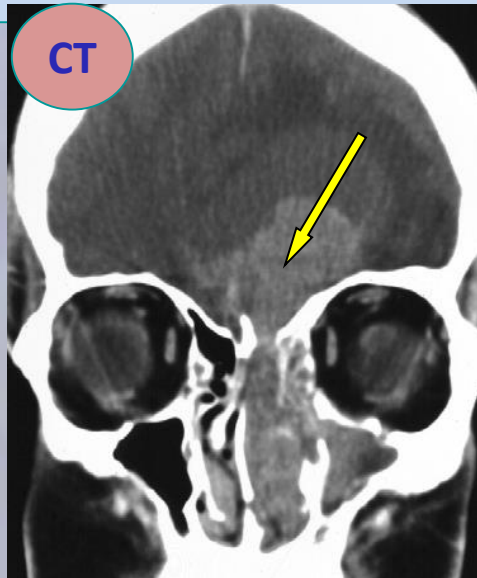


MR



CT – MR Acc: >90%

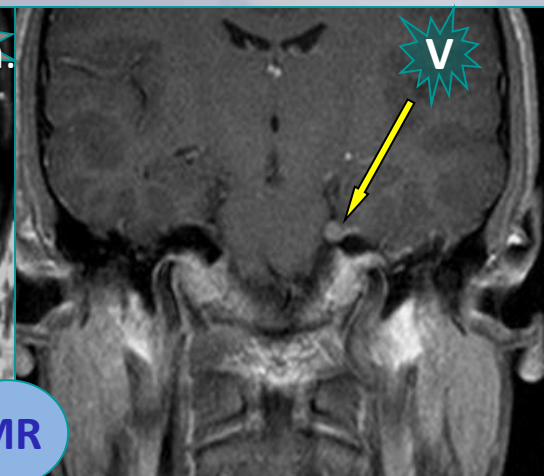
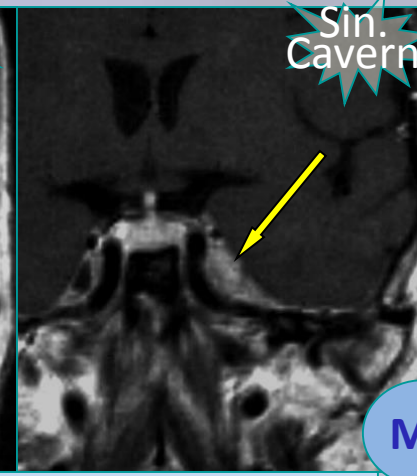
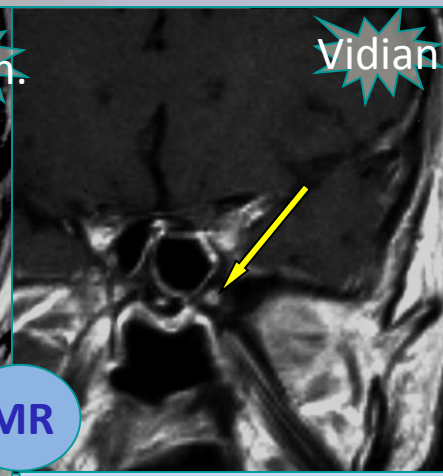
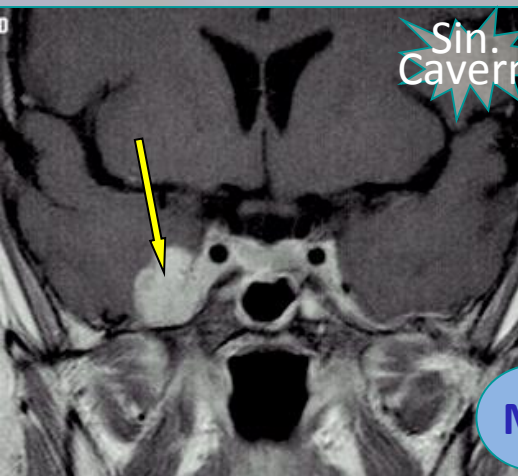
Intracranial TU extension - CT/ MR



epidural

dural

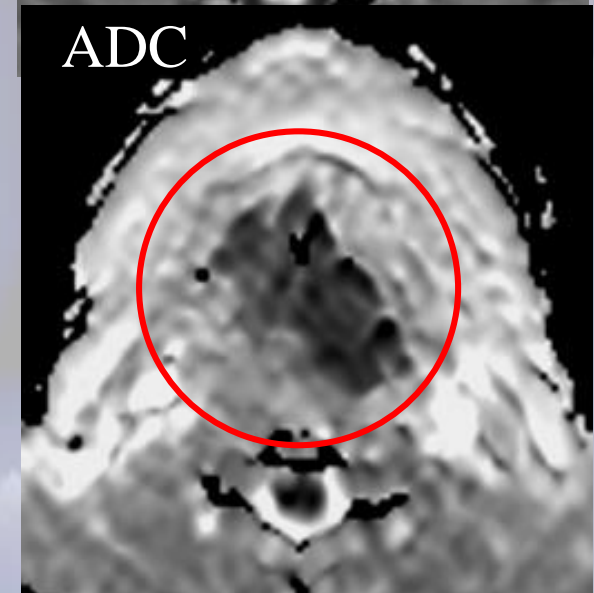
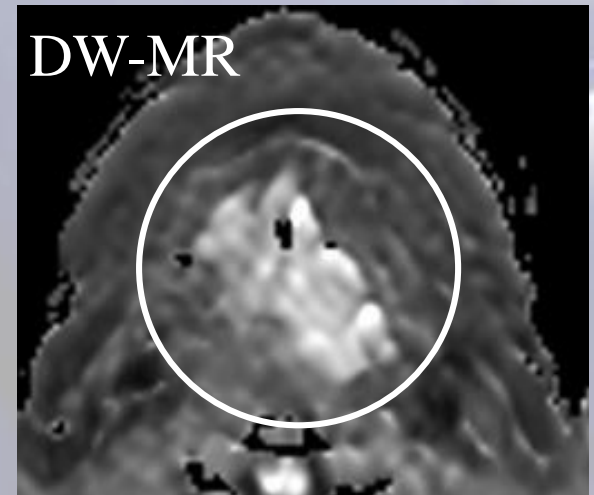
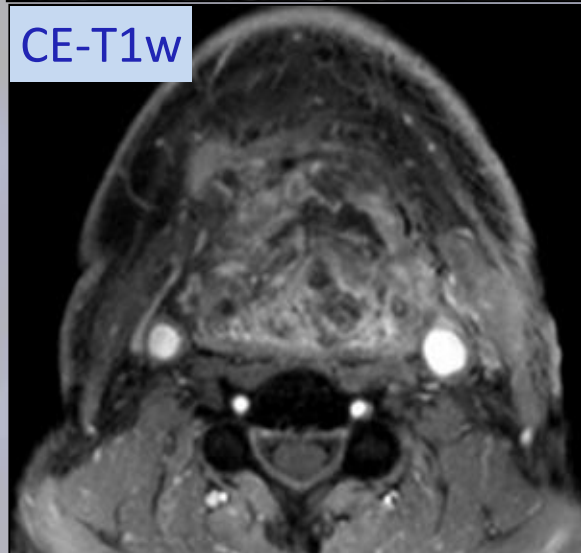
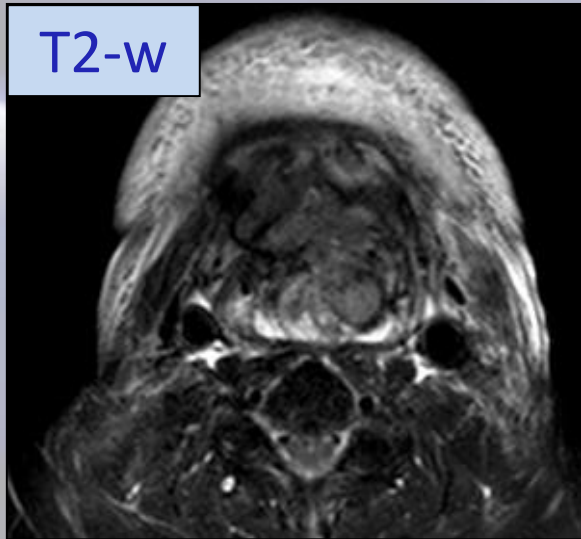
intracerebral



Perineural TU spread

Supraglottic residual carcinoma – MP-MRI

Restricted diffusion within the residual tumor tissue

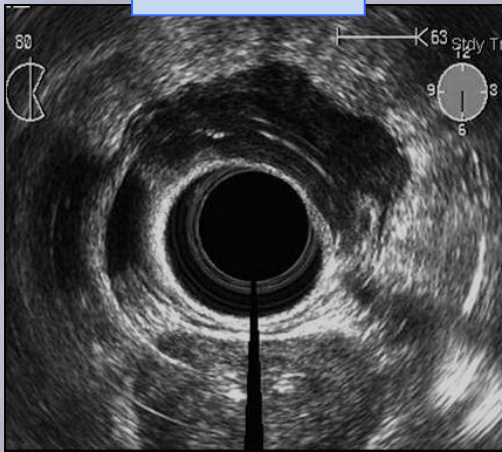


RECTAL TUMOR

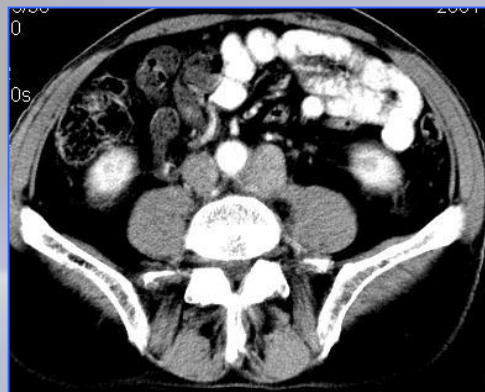
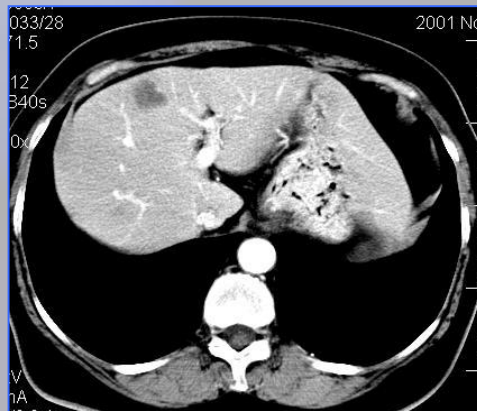
Multimodal Imaging

- **US** Transabdominal US for general abdominal information
Endorectal US – intramural TU extension
- **MP-MRI** best evaluation for tumor extension beyond the wall,
relation to the adjacent organs, lymph nodes, liver
- **CT** to evaluate advanced TU extension (thoraco-abdominal-pelvic)
- **US/CT guided biopsy** (liver)
- **PET/CT** whole body information – distant / recurrent Tu

EUS



CT
Rectal ca
Acc 70-85%



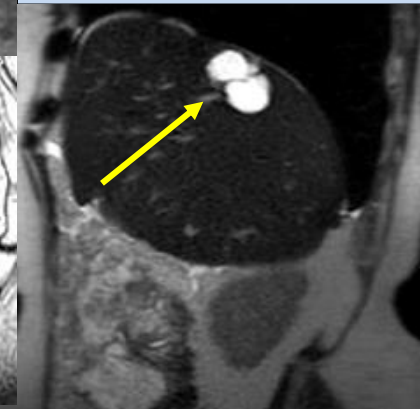
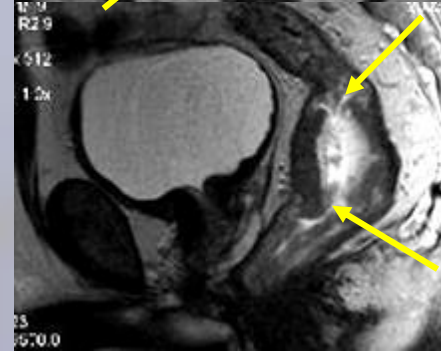
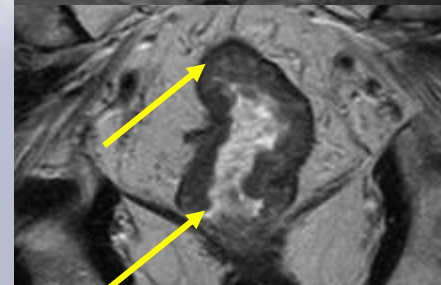
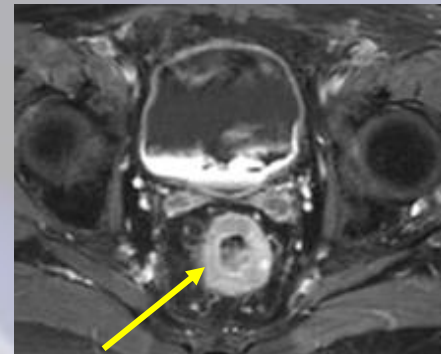
RECTAL Cancer

Clinical examination: rectoscopy

MRI

Acc>90%

Liver met
Acc>90%



Imaging in **PROSTATE** cancer

- ❖ **US** – general abdominal and pelvic informations
 - Transabdominal US
 - Endorectal US
 - Color- Doppler US
- ❖ **MP-MRI (T2-w, DW-, Dyn-MRI)**
 - Reliable pelvic status, T/N staging for prostate, recurrent tu?
- ❖ **Bone scan** – bone metastasis
- ❖ **CT**- to evaluate advanced Tu and dissemination
- ❖ **PET/CT** – for recurrent tumor or metastases

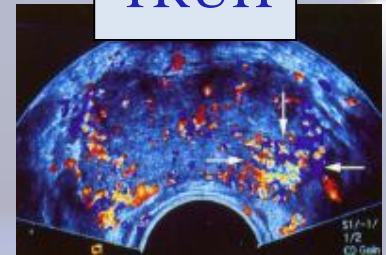
Prostate cancer

- **Clinical Dg:**
 - PSA – **not reliable for Dg. and staging**
 - Organ specific (*not tumor specific!*)
 - norm: >3-4-6 ng/ml (age dependent)
 - Good for follow-up
 - PCA – **better, but not sure**
 - DRE (**digital rectal exam**) – suspected tumor, usual underestimation
- **Sextant / TUR (transurethral) biopsy (core)**

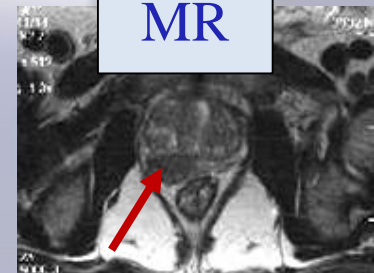
Imaging Dg:

- ❖ **TRUH** – orientation about the structure
guided biopsy with MRI image fusion
- ❖ **MP-MRI** – Dg, staging, therapy monitoring, recurrent tu?
- ❖ **Bone scan** – **bone metastases**
- ❖ **CT** – advanced Tu stages, TU **dissemination**
- ❖ guidance for radiotherapy
- ❖ **PET-CT:** recurrent tumor, therapy monitoring, recurrent Tu?

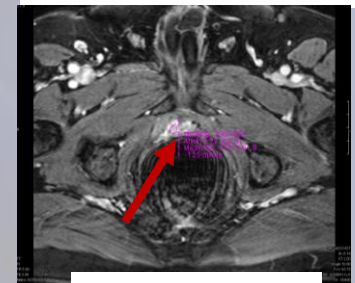
TRUH



MR



Tu broke through the capsule - T3a

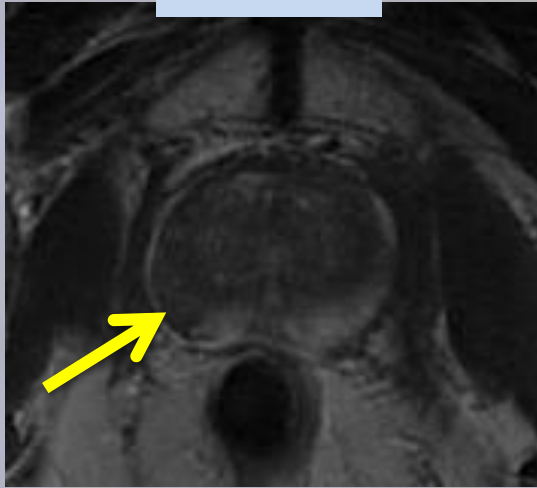


Recurrent Tu.

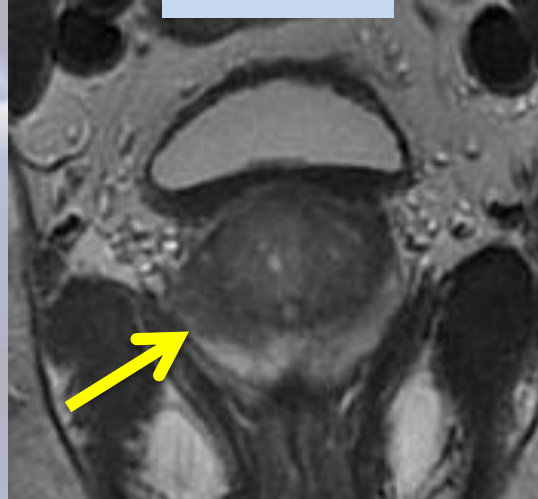
Prostate cancer

„T” Staging: MP-MRI (T2b)

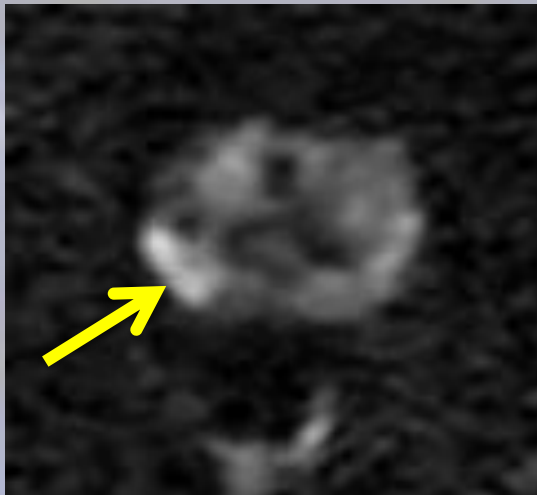
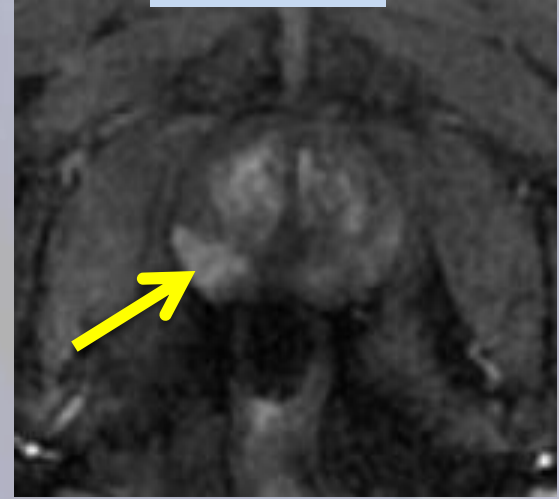
T2-w axi



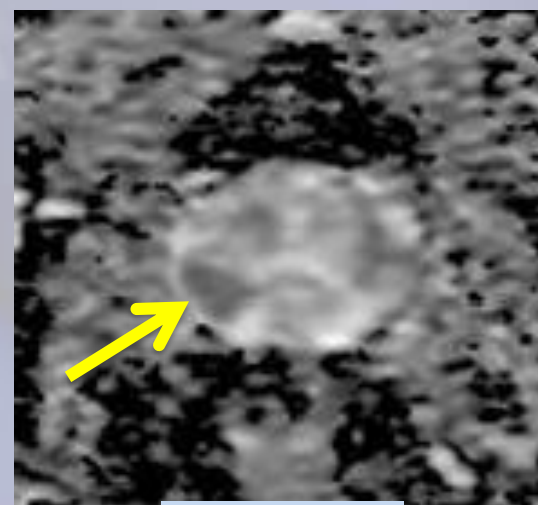
T2-w cor



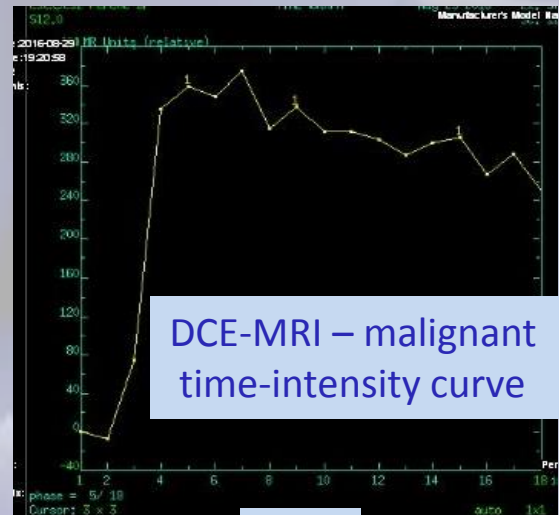
DCE-MRI



DW-MRI-b1000



ADC-MRI

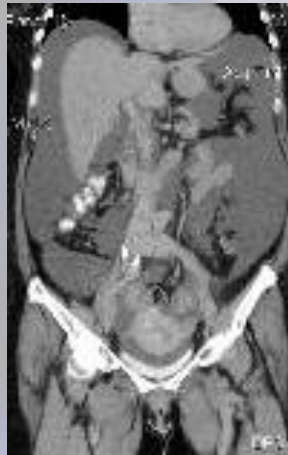


DCE-MRI – malignant time-intensity curve

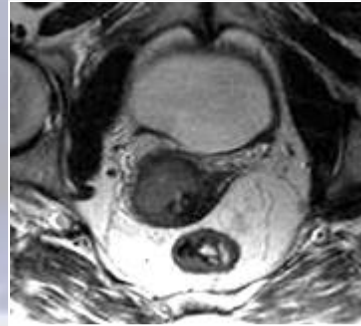
TIC

Imaging in gynecological tumors

- **US** – orientation
 - Transabdominal US (+Doppler)
 - Endovaginal US (+Doppler)
- **MRI – STAGING!**
- **CT** – advanced TU extension
OVARIAN!



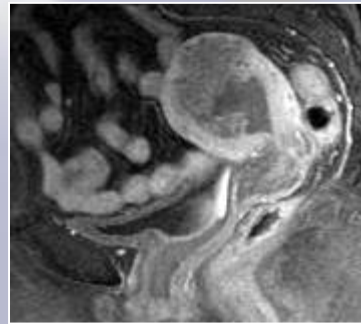
- **Guided /UH, CT/ biopsy**
- **PET/CT** – metastases, recurrent TU



Cervix ca.

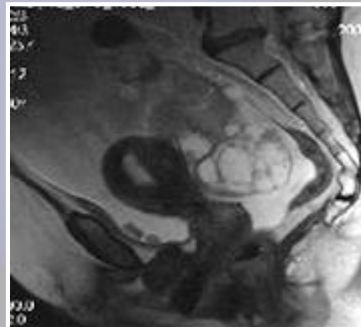
MR-ACC:>95%

Lymph node: 70-80%



Corpus tu.

MR-ACC:> 90%



Ovarian ca.

MR-ACC: 89-99%

Conclusion

- ❖ **CXR:** the evaluation of tumor is limited
- ❖ **US:** excellent for the evaluation of soft tissues, abdominal organs and excellent tool for tissue sampling, BUT don't forget the limitations!
- ❖ **MRI/CT** are basic modalities for cancer evaluation
 - ❖ CT & MR: complementing each other
 - ❖ Advantages of MR: better soft tissue resolution, multiplanar imaging, functional measurements
 - ❖ Advantages of CT: faster, less motion artifacts, whole body information, better assessment of cortical bone.
- ❖ **PET-CT:** detection of distant metastases, recurrent diseases, to evaluate therapy response

Optimal treatment is based on multidisciplinary decision

High quality imaging guides oncological treatment

Dyn-MR

CE-US

MD-CT

MD-CT

PET-CT

Interventions

DW-MR

Thank you for your attention!

...welcome in Blue Ball Street; Budapest

