Theoretical Bases of Clinical Medicine

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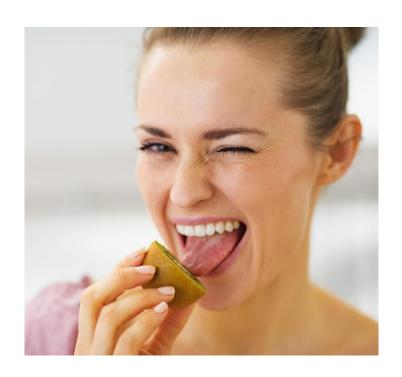






Aim of TZKM?

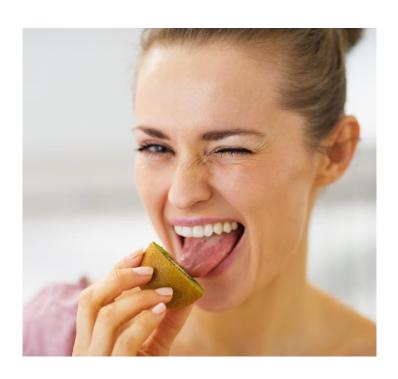
• "Taste" of clinical medicine...



Cíle předmětu TZKM?

• "Taste" of clinical medicine

- Breast Cancer
- Intravenous access in oncology



Questions to start with?

What to do if it the studies is too much?

- What is the most frequent malignancy in women in the Czech Republic?
- What screening programme is about to be started?
- Can a breathlessness (dyspnea) be a sign of breast cancer?
- What is a central venous access?
- How to treat a intravenous port (port-a-cath)?

Recommended literature...

- Mika Waltari: The Egyptian (1945)
- Richard Gordon: Doctor in the house (1952), Doctor at Sea (1953)
- Samuel Shem: *The House of God* (1978)



Medical students congress in ancient India (700 B.C.)

- Samudrah ivah gumbheeram
- naivah shakyam czikitsitam
- vaktum niravaasheshainah
- Schlokhanam ayutaïr apih



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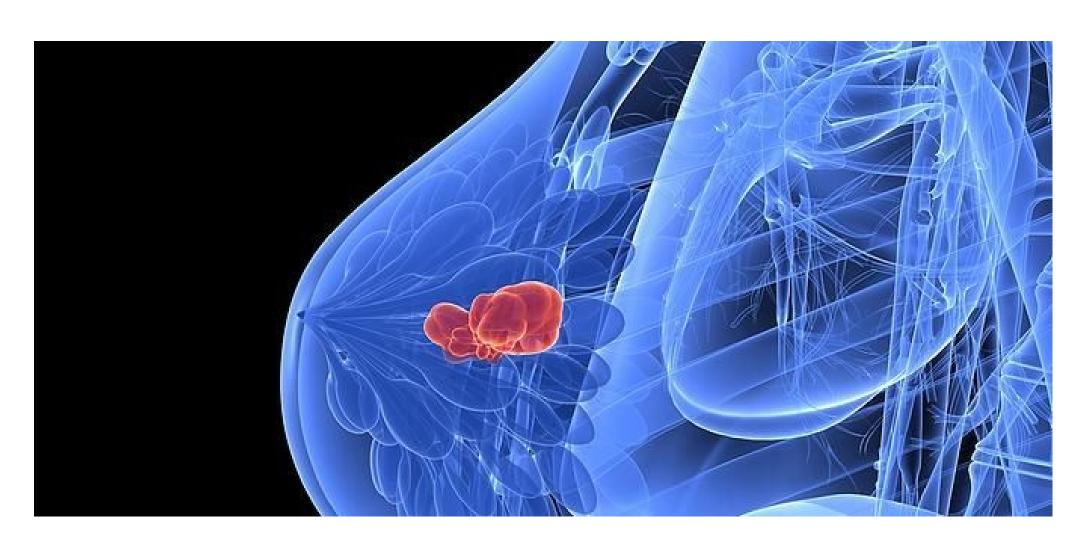
Medical students congress in ancient India (700 B.C.)

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- Uneasy is the Medicine
- Deep as a wide sea
- It is not possible to explain it in whole
- Not even in hundreds of thousands of verses
- (PhDr. Miltner- Lékařství staré Indie, Avicenum 1986)

Breast cancer



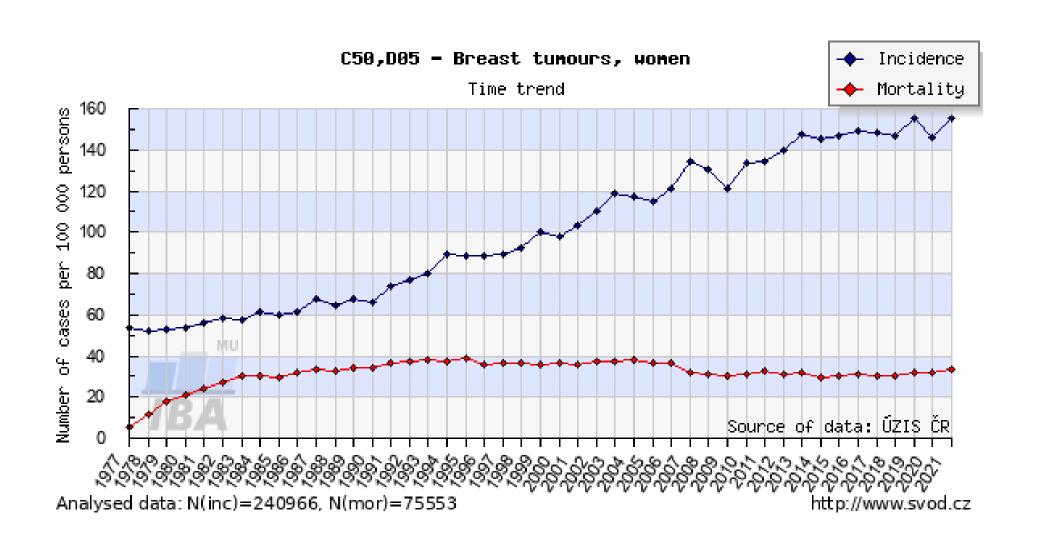
Incidence of the most frequent malignancies (CZE)?

order	males to 100k inhab.	to	females o 100k inhab	All to 100k inhab.	
1	?	?	•	?	
2					
3					
4					
5					

Most common malignancies- incidence Health statistics Institute: Neoplasms 2018

order	males to 100k inhab.		females to 100k inhab		All to 100k inhab.	
1	Skin nonmelanoma	287	Skin nonmela noma	250	Skin nonmelano ma	26 8
2	prostate	152	breast	133	colorectal	68
3	colorectal	83	colorectal	54	lung	61
4	lung	79	lung	43	kidney	29
5	kidney	38	Uterus (corp)	36		11

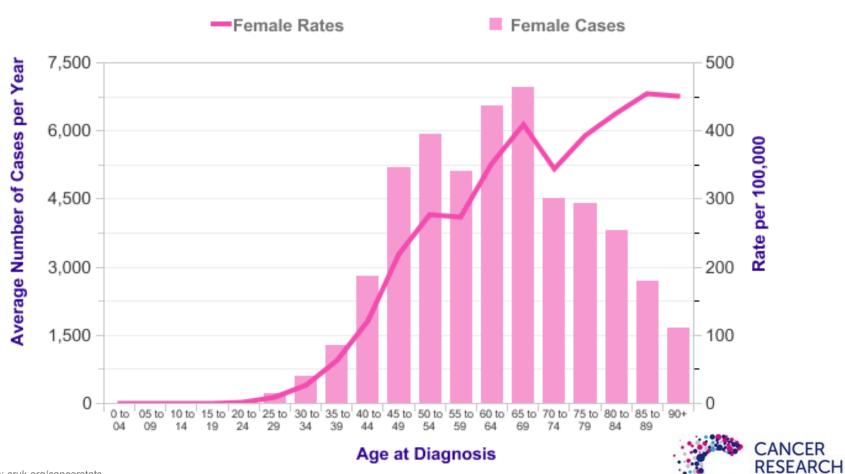
Incidence and mortality- Breast cancer CZE



Incidence BC age related

Breast Cancer (C50): 2011-2013

Average Number of New Cases Per Year and Age-Specific Incidence Rates per 100,000 Population, Females, UK





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Clinical signs What does a patient come with??

Clinical signs

- Breast resistance lump
 - Upper lateral quadrant most frequent
- Edema of skin
- Erythema of skin
- Retraction of skin, ulceration
- Inversion of mammila
- Painless afections
- General symptoms fatigue, weight loss, dyspnea...

Lump



Advanced tumour



Skin retraction + "mosquito bites"



Skin edema and erythema "peau d'orange"



Mammila inversion





Risk factors

Can I do something not to have it in future?

Risk factors

• Family history: breast cancer of 1st degree relative (parents, siblings, children)

• One - relative risk 1,5 - 2,0

• Two relatives 5,0

- Early period onset: before 12 yoa
- Late menopause: after 55 yoa
- Nullipara (no birth) + no breast feeding
- Combined hormonal substitution
- Smoking, low physical activity, night shifts?
- Benign breast afections: Atypical ductal hyperplasia
- Genetic factors, various 5-10 % of breast tumours

Genetic risk - example

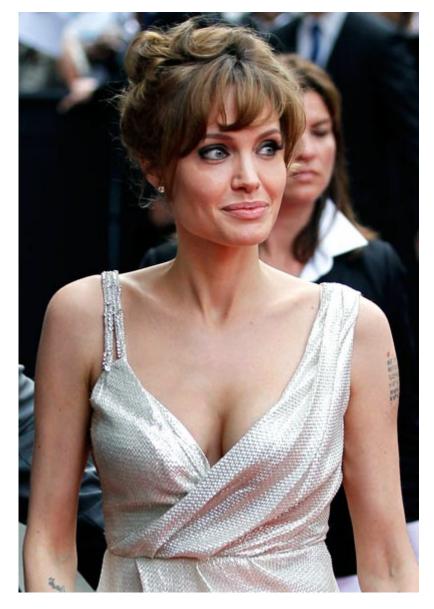
- BRCA1 a BRCA2 genes DNA repair homologeous recombination
- Risk of ca breast in **BRCA1** mutation = 80% (lifetime), ovarian ca 60%, **BRCA2** mutation 70%, resp. 25%
- Only prevention bilateral mastectomy + salphingooophorectomy

Likelihood of a 25-year-old woman surviving to age 70

(without screening or medical interventions to prevent cancer)

Group	Percentage surviving to age 70
BRCA1 mutation	53
BRCA2 mutation	71
Typical woman	84





Screening – general principles

- Cheap
- Nonivasive
- Highly sensitive (few patients "escape")
- Effective (survival benefit...)

• Doesn't have to be specific

Screening -oncology

- Gynecologic
 - Cytology smears from cervix
 - From onset of regular gyno assessment
- Mammar
 - Mammograph after 45yoa every 2 yrs
- Colorectal
 - Occult stool bleeding after 50 yoa (hidden!! bleeding) or
 - Scr colonoscopy after 55 yoa
- Lung ca low dose CT in risky population (start 2022)





- Soon:
 - Ca prostate? discussed

Diagnostics

How do we find it??







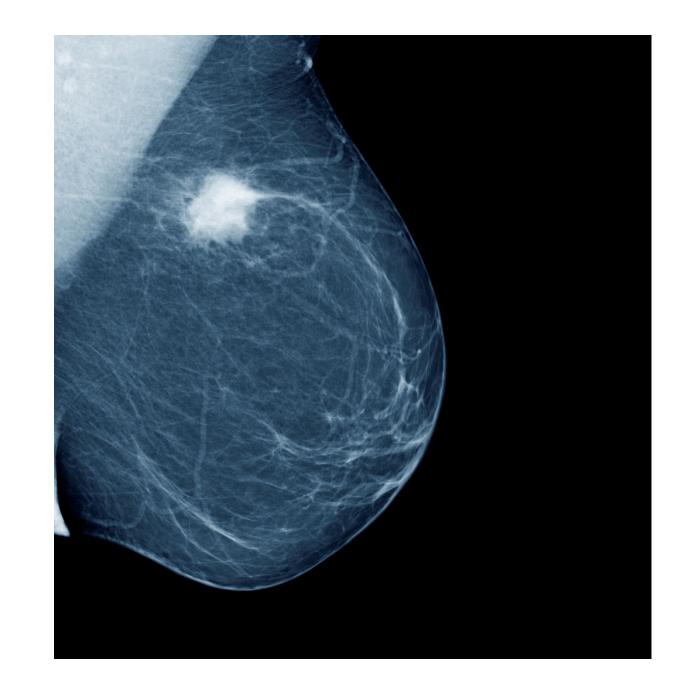
Inicial diagnostics

- 62-yo woman
- New lump left breast (upper lateral quadrant)
- Overall healthy and good shape

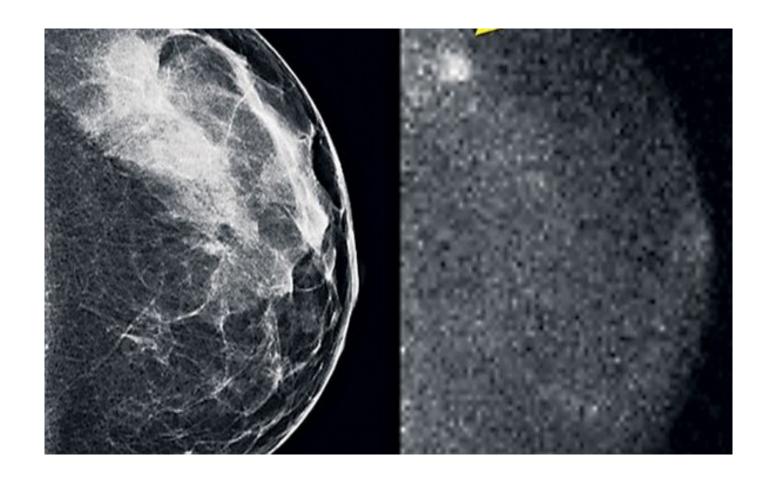
- What investigations???
- TNM staging...

Mammography

- Screening and diagnostic method for ca breast
- Very sensitive and specific
- Cheap and safe (low radiation)
- Size evaluation (in mm)
 - T stage
- Diagnostics of regional lymph nodes
 - N stage
- Sometimes US better or MRI



MMG, US

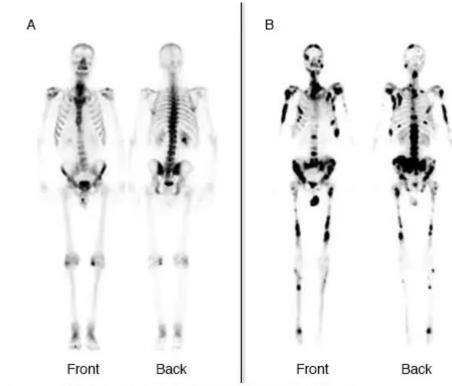


Staging Ca breast

T classification	Size, characteristics
ТО	No evidence of primary tumour
T1	Tumour ≤20 mm largest diameter
T1a	Tumour >1 mm but ≤5 mm largest diameter
T1b	Tumour >5 mm but ≤10 mm largest diameter
T1c	Tumour >10 mm but ≤20 mm largest diameter
T2	Tumour >20 mm but ≤50 mm largest diameter
Т3	Tumour >50 mm v největším rozměru.
T4	Tumour whatever size infiltrating chest wall and/or skin (ulceration, skin lesions)
T4a	Infiltrating chest wall (not only the muscle)
T4b	Ulceration and/or edema (including peau d'orange) of skin
T4c	either T4a or T4b.
T4d	Inflammatory carcinoma

Distant metastases - staging

- M stage
- Assessment:
 - Chest: Xray, CT scan
 - Abdomen and pelvis: ultrasound, CT scan
 - Whole body PET/CT or PET/MRI, wb CT
 - Bones scintigraphy
 - Brain MRI or CT (with contrast!!)



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Staging and survival – ca breast

Stage	Т	N	M
1	T1	NO	MO
IIA	T0	N1	MO
	T1	N1	MO
	T2	N0	MO
IIB	T2	N1	M0
	T3	NO	MO
IIIA	T1-3	N2	MO
	Т3	N1	MO
IIIB	T4	Any N	MO
IIIC	whatever T	N3	MO
IV	whatever T	whatever N	M1

Stage	5-y OS		
I	95%		
IIA	85%		
IIB	80%		
IIIA	67%		
IIIB	41%		
IIIC	49%		
IV	15%		

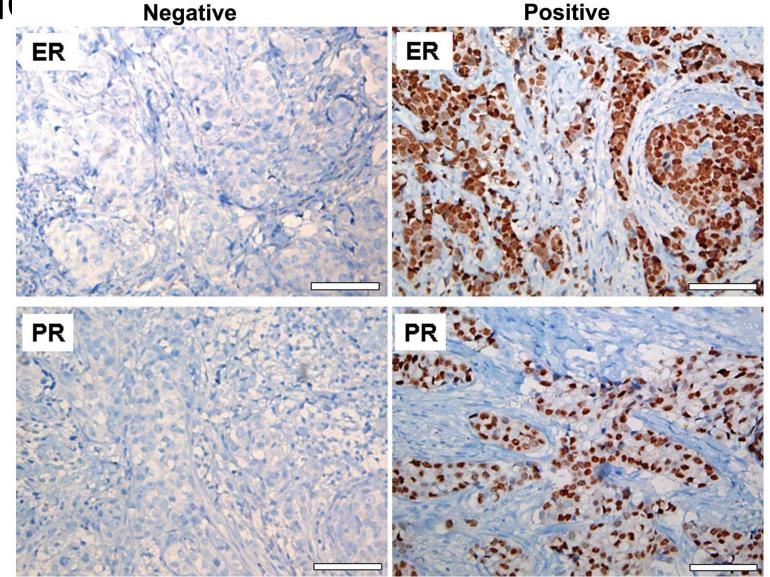
Histology

what do we deal with??

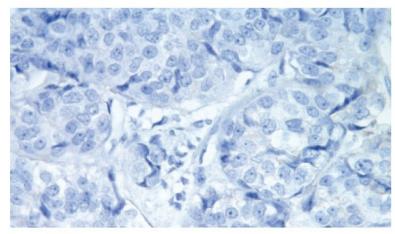
Histology = cells and stroma

- Morphology "typing":
 - Ductal (85%), lobular, medullar, mixed
- Grade of differentiation "grading"
 - Grade 1 well diferentiated (good prognosis)- grade 4 undifferentiated (poor)
- Morpho-biology:
 - Receptor expression:
 - Estrogene
 - Progesterone
 - HER2 receptor
 - **Ki-67** marker of proliferation (%)

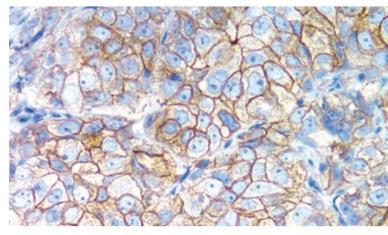
Expression ER a PgR - immunchist Regative



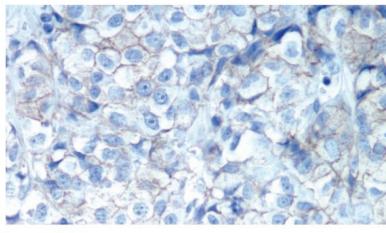
HER2 expression - immunohistochemistry



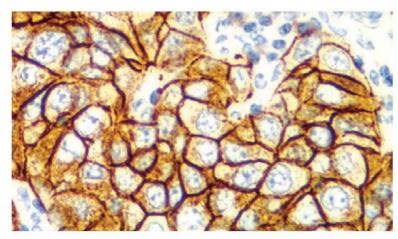
Score: 0 (40x)



Score: 2+ (40x)



Score: 1+ (40x)



Score: **3+** (40x)

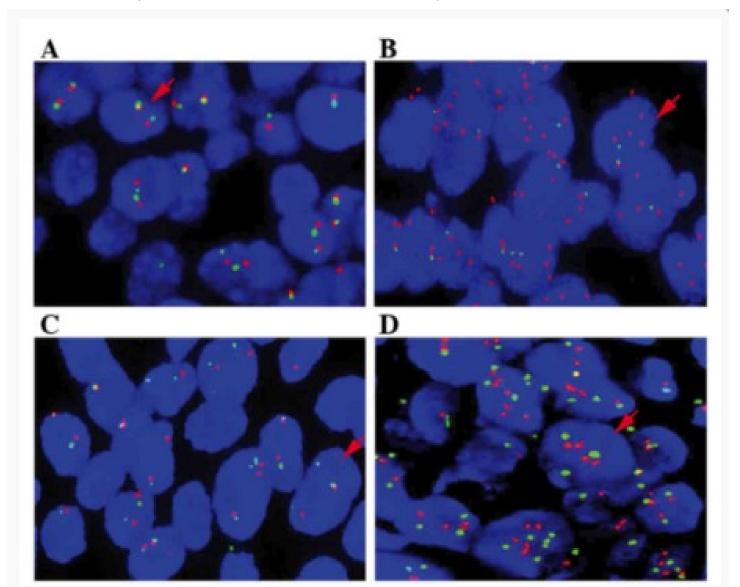
• A+C:

• No amplification HER2

B+D:

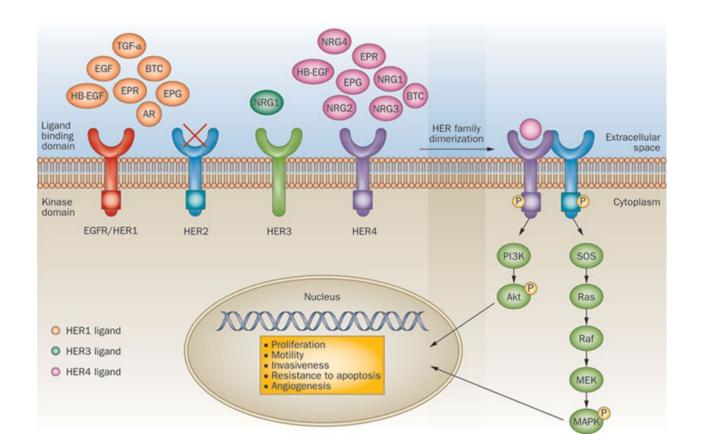
amplification

FISH



HER2 positive BC

- HER2 transmembrane receptor
- Active through homodimerization (HER2-HER2) or heterodimerization HER2-HER3 or HER2-EGFR



Triple negative breast cancer -TNBC

ER negat, PR negat, HER2 negat



Only modality – chemotherapy

- not hormones
- not targeted (perhaps besides-VEGF therapy)

Immunotherapy??

Subtypes breast cancer

Luminal A

ER+, PgR+, HER2-, Ki67 low

Luminal B

ER+, HER2+ or -, and another risk factor: PgR negative, Ki67 high

Triple negative

ER-, PR-, HER2 negat

HER2 overexpression (amplified)

Subtypes breast cancer

Median overall survival

```
OS (months)
                                 site of metastases

    luminal A

                  26,4
                                   bones, liver

    luminal B

                   19,2
                                   bones, liver, lungs
• luminal/HER2+ 15,6
                                   bones, liver, brain
• HER2+
                                  bones, liver, lungs, brain
                    8,4

    Triple-negative

                  6,0
                                  lungs, brain
                 (p < 0.001)
```

Therapy

how to cure?

...if not cure, how to prolong life?

Principles of therapy

- Localized disease- attempt to cure
 - **Resection of primary tumour** only possible curative approach
 - **Neoadjuvant** therapy in some cases before the operation chemotherapy or hormones
 - Aim is to shrink (downstage) the tumour and lessen the extent of surgery
 - Adjuvant therapy after the operation
 - Aim to lower the risk for relapse
 - Toxicity (temporary) is not too relevant
- Metastatic disease we can prolong life
 - **Systemic** treatment chemotherapy, hormones, targeted treatment
 - Toxicity relevant! Quality of life!

Operation

remove it, if we can!!!

Mastektomy

• Total mastektomy – removal of breast in whole

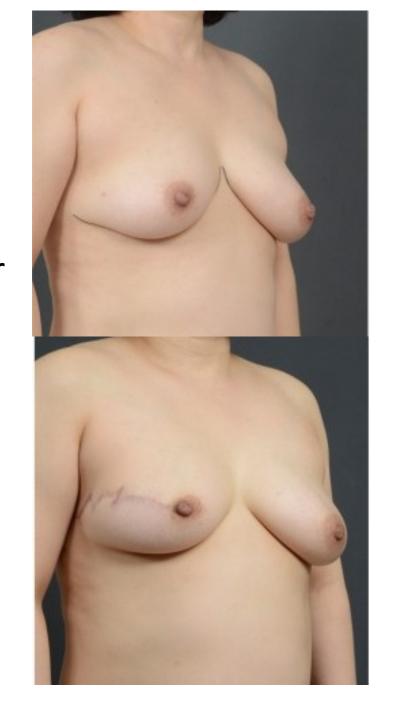




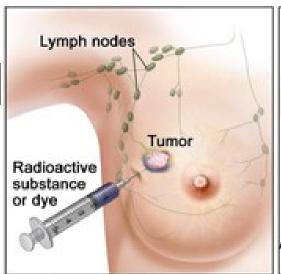
Partial mastektomy

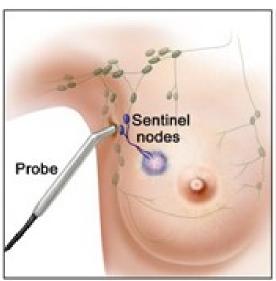
 Breast saving operation – removing the tumour and small amount of surrounding tissue

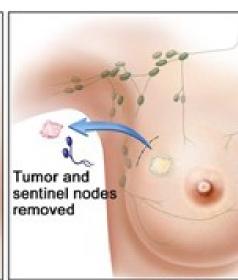
Aim to save natural shape and form of the breast



Operation in axil







Axillar dissektion - ALND

- Incision in axilla, removal of 10–40 lymph nodes of level I and II
- All patients with lymf nodes involvement ("positive")
- Real risk of long term lymfedema

Sentinel node biopsy – SNB

- Sentinel lymph nodes first to be infiltrated (first to pass lympha from tissue surrounding the tumour
- Removal of 1-3 lymph nodes in all patients with NO signs of LN involvement
- I positive (infiltrated), ALND is pursued

Risks of nursing on an arm after ALND

- infection (i.g. erysipelas)
- cellulitis
- Lymphedema or its progression
- The risk is relative
 - e.g. Hand surgery can be safe
- Recomm.: choose the other arm if possible for blood taking, BP measurement
 - Not aplicable for emergency!!! (risk/benefit)

Adjuvant treatment

prevent relapse!!



Adjuvant treatment

- After the operation
- To minimize risk of relapse
 - Aim to kill residual microscopic disease
- Multimodal treatment:
 - Chemotherapy 4 months
 - Targeted treatment (HER2 posit.) 1 year
 - Radiotherapy 5 weeks
 - Hormones (ER/PgR posit.)— 5–10 years or more

Chemotherapy - whom for?

- Selected patients with risky tumours:
 - Large tumour
 - Positive lymph nodes
 - Biologically aggressive disease triple-negative, HER2 positive

Chemotherapy and breast cancer

Most used cytotoxic drugs:

- Antracyclines Doxorubicin, Epirubicin
- Taxanes Paclitaxel, Docetaxel
- Cyclophosphamide, 5-Fluorouracil only in combinations

Combinations in adjuvant setting:

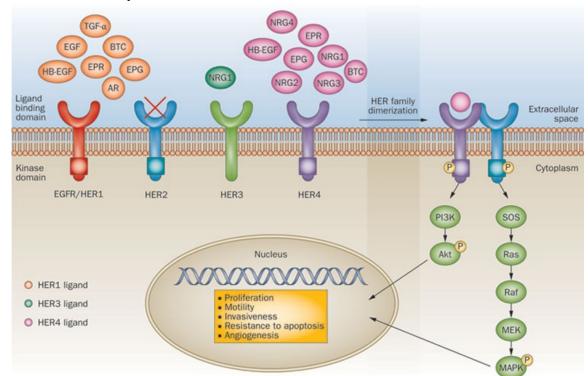
- AC doxorubicin + cyclophosphamide
- AC followed by paclitaxel
- FAC flurouracil + doxorubicin + cyclophosphamide
- FEC flurouracil + epirubicin + cyclophosphamide
- TAC docetaxel + doxorubicin + cyclophosphamide
- CMF cyclophosphamide + methotrexate + 5-fluorouracil

Metastatic setting:

 Combination (more effective) or monotherapy (more gentle) – paclitaxel, epirubicin, vinorelbin, capecitabine

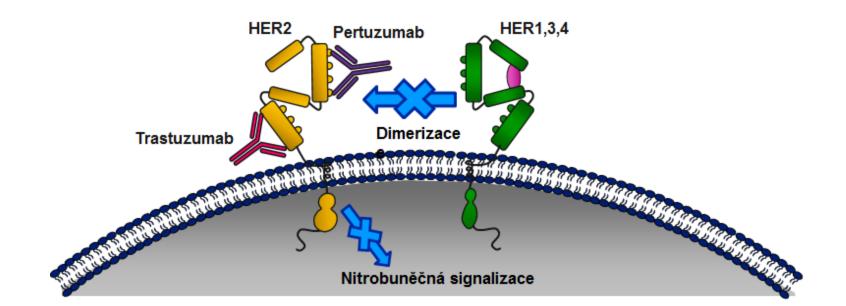
Targeted treatment

- Used in **HER2** positive breast cancers, approx. **15% pts**
- HER2 transmembrane receptor, EGFR family (HER2=EGFR2)
- Activated by homodimerization (HER2-HER2) or heterodimerization (HER2-HER3 nebo HER2-EGFR)



HER2 positive breast cancer

- anti-HER2 therapy monoclonal antibodies against HER2 receptor
- First used **Trastuzumab** (Herceptin™)
- Later generations pertuzumab, T-DM1
- Adding to chemo adds effectléčby



Hormonal therapy

 Hormonal receptors expressed in 70 % BC (estrogene or progesterone receptor) = hormone sensitive tumour

Usually low or moderate aggressive tumours (Luminal A a B)

- Sensitive to hormonal treatment:
 - Tamoxifen (Selective Estrogene Receptor Modulator SERM)
 - Aromatase Inhibitors
 - Non-steroidal AI (anastrozole, letrozole)
 - Steroidal (exemestane)
 - Direct ER inhibitor (fulvestrant)

Hormonal therapy- mechanism of action (MOA)

1. competition – on estrogene receptor, modulation of ER – tamoxifen, direct ER inhibitor - fulvestrant

- 2. inhibition blockade of sythesis of estrogene (inhibition of aromatase enzyme in fatty tissue) AI: anastrozole, letrozole, exemestan
- 3. ablation ovarian estrogene suppresion (farmacologic castration LH-RH analogues goserelin, buserelin, leuprorelin, triptorelin)
- 4. adition adding estrogenes, androgens or gestagenes more in history

Hormonal treatment

- In adjuvant setting administered for 5-10 years
- Very effective in Luminal A subtype of BC
- Some patients can be saved from chemotherapy and use hormones only
- Treatment with low/minimal toxicity (heat flushes, bones and joints pain, artificial menopause, endometrial carcinoma and tamoxifen)

Radiotherapy

- After operaci a chemotherapy
- Always follows partial mastectomy, sometimes total mastectomy (large tumour, positive LN)
- 5-6 weeks, dose 50-60 Gy
- Reduces risk of local relapse and improves OS
- Toxicity: dermatitis, skin deskvamation



Therapie of a disseminated incurable disease

no cure, rather prolonging life and improving/keeping its quality

Terms and Definition

- Curative- aim to cure
- Palliativní- (noncurative), cure not expected
 - Aim to prolong survival, quality of life
 - Invaziveness of the procedures according to life expectancy and pts' wish (years?? vs. days??)
- Causal treatment (anticancer)
- Symptomatic treatment (symptoms)

Therapy of metastatic disease

- SR=ER/PgR positive + breast cancer (Luminal A/B subtypes) -
 - hormonotherapy very effective
 - Tamoxifen \rightarrow Al \rightarrow Fulvestrant

• If not effective or SR- chemotherapy

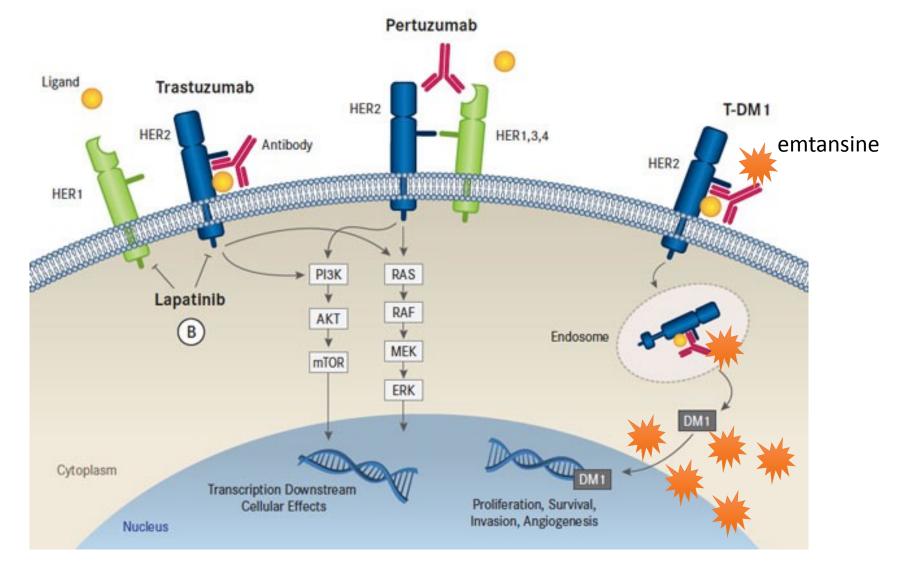
• HER2 positive tumours- combination with **targeted treatment** (trastuzumab, pertuzumab, T-DM1)

Chemotherapie

- Monotherapy better???
- If progression/toxicity, followed by further lines
- Antracyclines:
 - Doxorubicin
 - Epirubicin
 - liposomal doxorubicin
- Taxanes
 - Paclitaxel
 - Docetaxel
 - Nab-paclitaxel
- Vinca alkaloids
 - Vinorelbine

- Antimetabolites
 - Capecitabine
 - Gemcitabine
 - Fluorouracil
- Platinum derivates
 - Carboplatin
 - Cisplatin
- Other
 - Cyclophosphamide
 - Methotrexate
 - Eribuline

Anti-HER2 therapy



Disseminated hormone-dependent ca breast

- Hormonotherapy
 - Preferred
 - Chemotherapy
 - If quick response and regression is needed
- Response anticipated within
 - Homones 2-3 months
 - chemotherapy 2-3 weeks

Specific and supportive care

- Bone mets bone-modifying agents (BMA) bisphosphonates, denosumab
- Painful bone mets radiotherapy
- Brain involvement surgery, radiosurgery, radiotherapy
- Pleural effusion drainage, talcage

- Supportive care- anaemia, pain, nausea, neuropathy, nutrition, intravascular access management...
- Psychologic, psychosocial and spiritual help

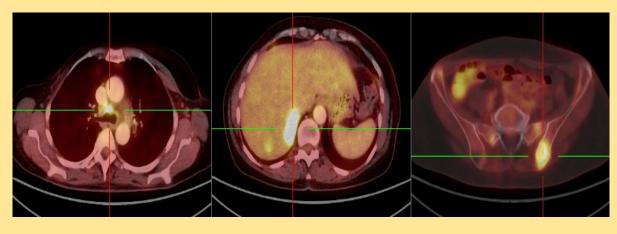
Case: Patient – 62yo woman

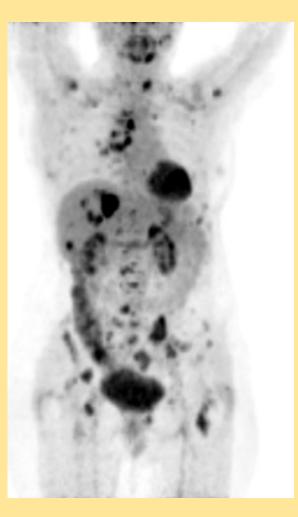
- 1999- Carcinoma mammae I.dx- pT1 pN1 (1/12) M0
- Low differentiation carcinoma ER-, PR+, G3
- St.p. RAME, adjuvant CHT 6x FAC and 5 yrs adjuvant hormones Tamoxifen
- 9/2011- relapse- soft tissues, bone mets, liver, tumor marker elevation
- Biopsy from liver lesions, phenotype: ER 80% PR 70%, Ki 67 25%

Case-continued

- 1/2012 to 11/2012 paclitaxel 1x weekly effect: minimal response
- 11/2012 to 1/2013 capecitabine effect of progression
- 1/2013 XENA (capecitabine+vinorelbine) to 4/2013, effect- progression (OSS, HEP)
- Still good shape, performance status (PS) 0 no limitations, working, active

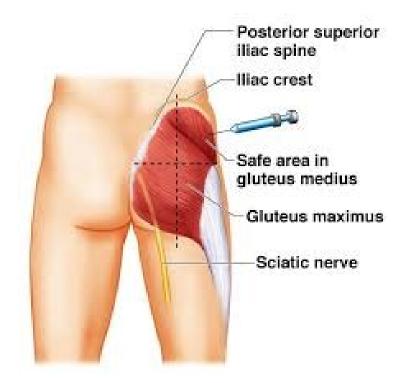
Case continued, PET/CT 4/2013 (after chemo)





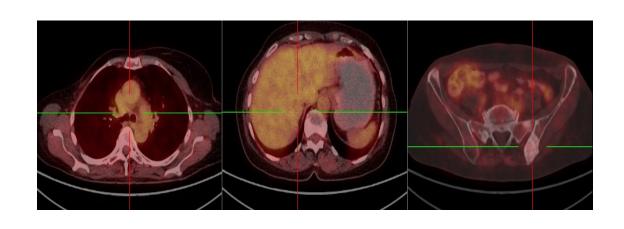
Case continued

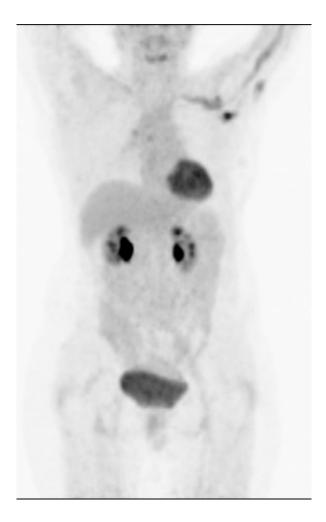
- hormonotherapy fulvestrant
 - 500 mg every 4wks intra muscular





Within 10mths effect of: overall improvement, all metastases regression (PET/CT - 3/2014)





Case 2: Pt, 52 years old

- 2/2010- ca left breast pT1b pN0 MO
- ER 80% PR 80%, low proliferation, HER 2 negative
- Parcial mastektomy + SNB, radiotherapy, adjuvant Tamoxifen
- 4/2014 dg liver lesions

Case 2: continued



Case 2: continued

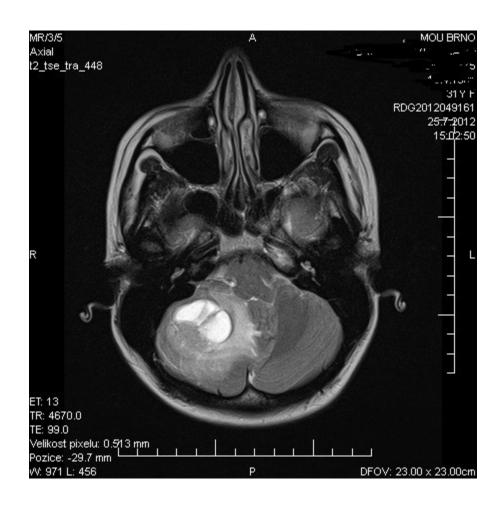
- 5/2014 biopsy liver lesions
- Histology: lymfocytes infiltrating liver

- Radiologist: typical metastases CT, US
- 7/2014- repeated biopsy liver-
 - Again- no cancer

Case 3: Female 29yo

- 2/2012 dg invasive ductal carcinoma
- triple negative, high proliferation (Ki-67 90%)- rapid growth T2N1M0
- Neoadjuvant chemo- AC (doxorubicin +cyclophosphamide)- effect after 3 cyclespropo 3 cyklech – progrese
- Změna na docetaxel- progrese after 2 cycles, carboplatin added- progression
- Mastectomy + ALND
- 9/2012-brain metastases

Case 3: Female 29yo



Ca breast: Summary

- Frequent malignancy in women
- Not one dissease various subtypes (biology, behaviour)

 Complex multimodal treatment- operation, radiotherapy, chemotherapy, hormones, targeted treatment

Supportive care complex and important

Side effects of chemotherapy

- Blood cell production (bone marrow, hematotoxicity)
- Germinal cells (spermias, eggs)
- Mucose (GI tract et al.)
- Nerves (feeling, movement, hearing)
- Organs heart, kidneys, liver, brain

Often irreversible, long term

Pří

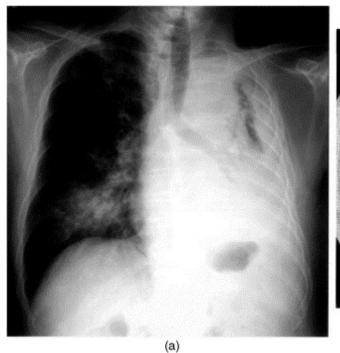






Nežádoucí účinky cílené léčby

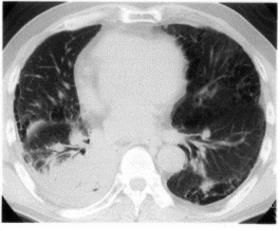
- Alergická reakce
- Únava
- Vyrážka/jiná kožní toxicita
- Průjmy
- Hypertenze (vysoký krevní tlak)
- Proteinurie (bílkovina v moči)
- Další (útlum krvetvorby, krvácení, nechutenství, hubnutí, chřipkové příznaky ad.)
- Většina reverzibilní (po vysazení léku se upraví)
- Některé vedou k přerušení/vysazení léčby
- Irreverzibilní také
 - Poškození orgánů: (srdce snížení funkce, plíce vazivovatění, snížení funkce, neinfekční záněty)





Pulmonary fibrosis





83

Skin symptoms – acne, blisters



Treatment of rash effective

A.



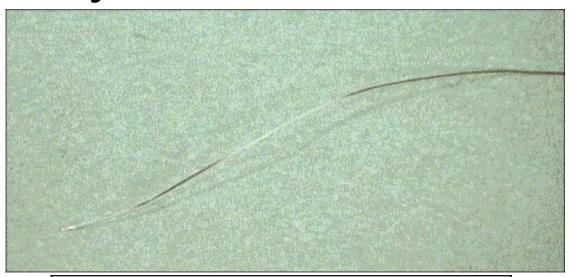
B.

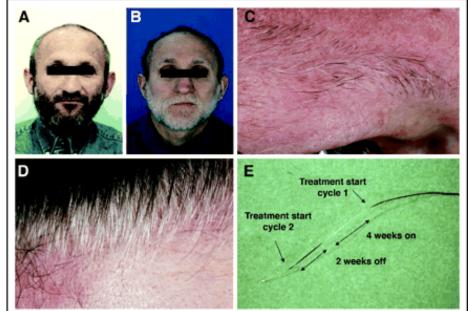


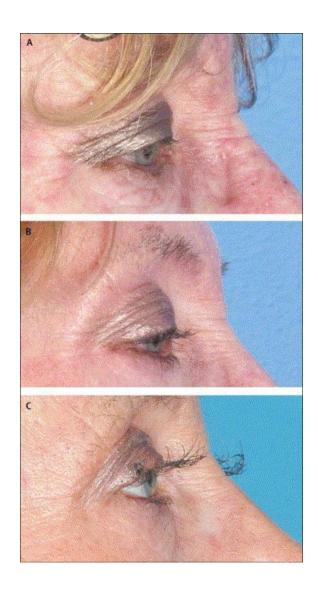




Hair + eye lashes







Vascular /intravenous access in oncology

- Complex
- Dynamic evolution (material, methods)
- evidence based

- Professional society
 - Society for ports and permanent catheters
 www.sppk.eu
 - World Congress of Vascular Access (WoCoVA.com)
 - guidelines





What is right (state of the art)? - 2023

- Choose a proper access, right indication
- Right procedure (implantation)
- Check the position of the tip of catheter
- Complication- special and skilled personnel
- Interdisciplinary team?



Right function, minimal burden for the patient

Why interdisciplinary?

- Indication (what for?)..... oncologist
- Implantation.....surgeon, radiologist, oncologist
- Nursingregistered nurse

- complications..... surgeon, hematologist, etc.?
- state of the art? news? Legal aspects?

Everybody knows his part...

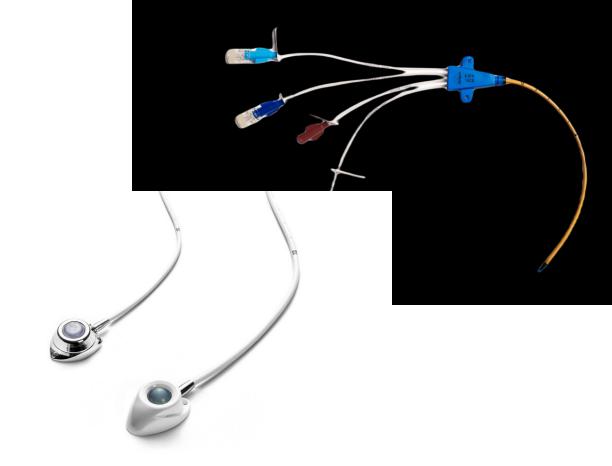
What is important to choose well?

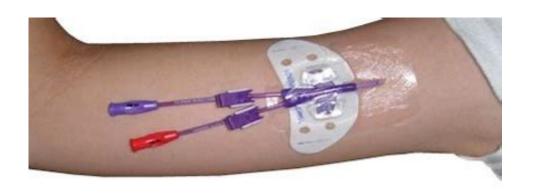
- For how long?
- What for? (infusions, chemicals, nutrition, blood taking)
- What will be administered? (pH, irritants, osmolality?)
- Where will be administered (hospital, home setting)
- Safety while inserting
- Infectious and thrombotic risk management
- Patient's preference (arm, chest)
- ...economics?

Cathegories

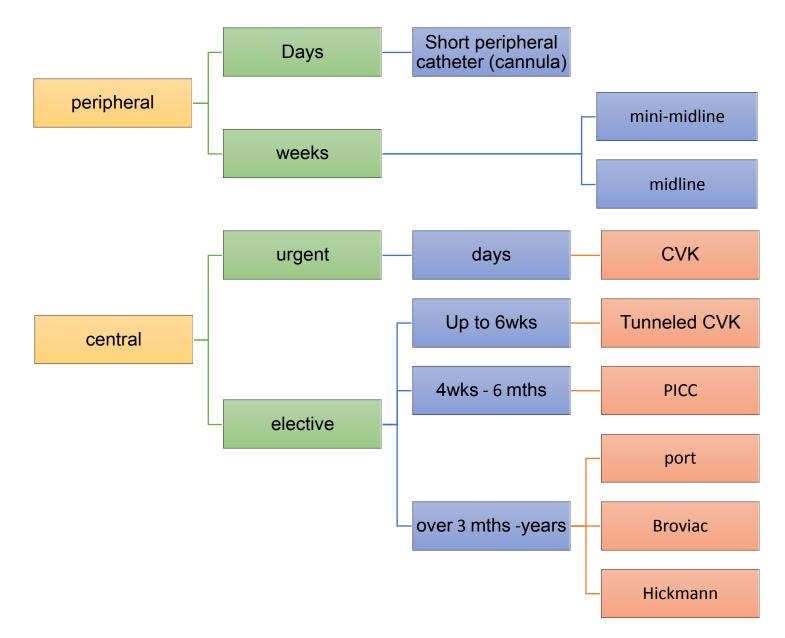
- Location of the tip
 - periferal
 - central

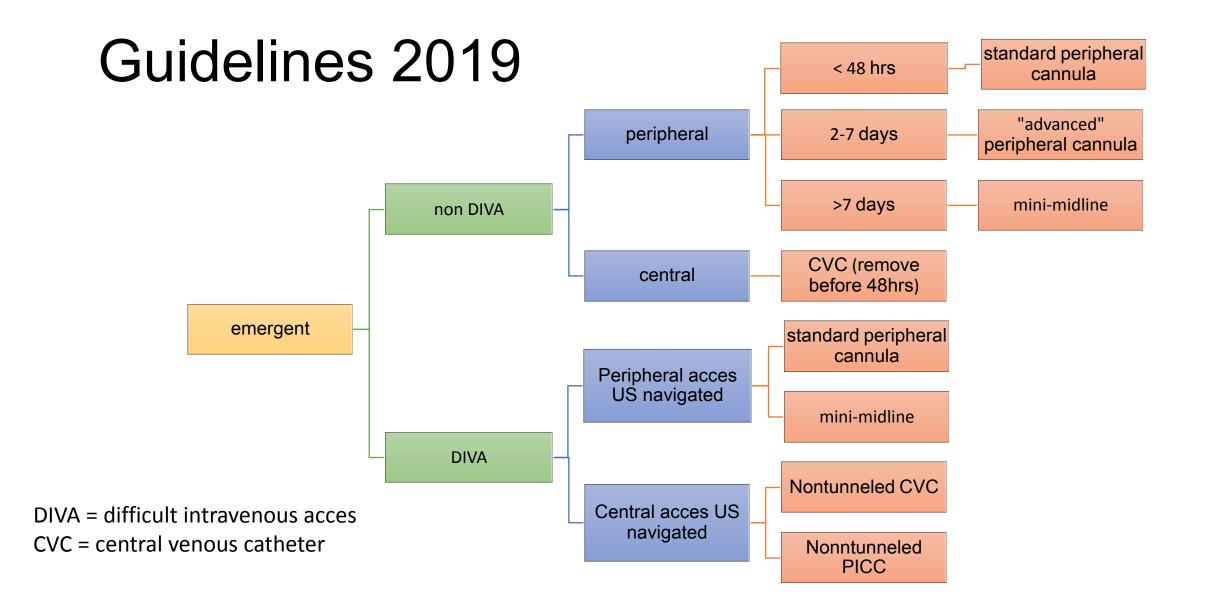
- Lenght of use
 - Short term (up to 7 days)
 - Middle term (ap cca 4-6 týdnů)
 - Long term = permanent (months even years)





Choice





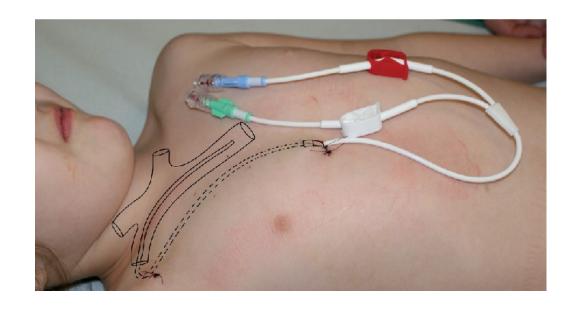
According to Italian guideline GAVeCeLT 2019 (Gli Accessi Venosi Centrali a Lungo Termine)



Purpose of tunnelization

• Exit site of catheter different from insertion site

- Why?
 - Optimal fixation
 - Easier nursing
 - Smaller risk of complication (infection)



ZIM = "Zone Insertion Method"

Robem Dawson - optimal choice of catheter exit site for PICC

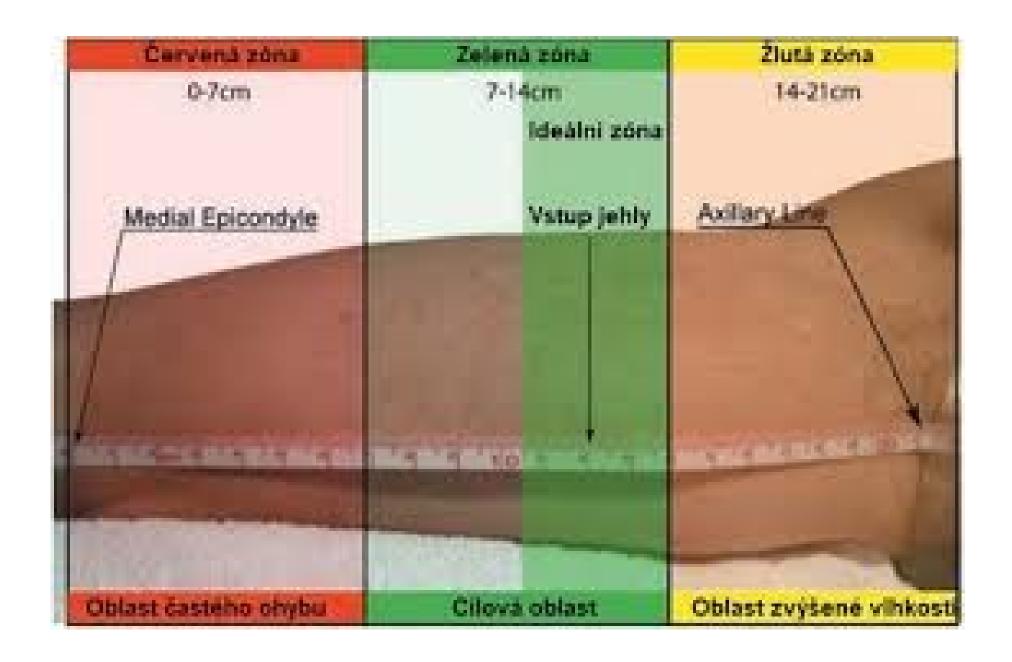
JAVA, 2011

PICC Zone Insertion Method™ (ZIM™): A Systematic Approach to Determine the Ideal Insertion Site for PICCs in the Upper Arm

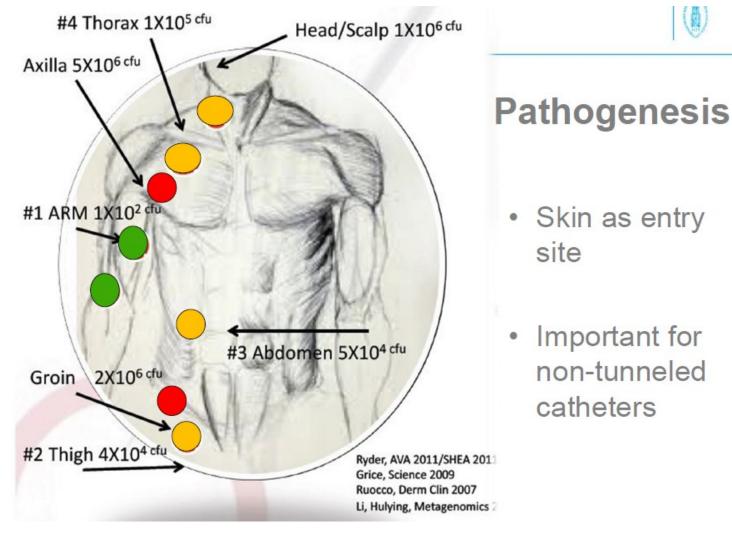
Robert B. Dawson MSA, BSN, RN, CRNI, CPUI, VA-BC

ZONE INSERTION METHOD (ZIM) Red Zone O-7cm 7-14cm Ideal Zone Needle Insertion Axist of Flexion Taiget Area Area of Moisture

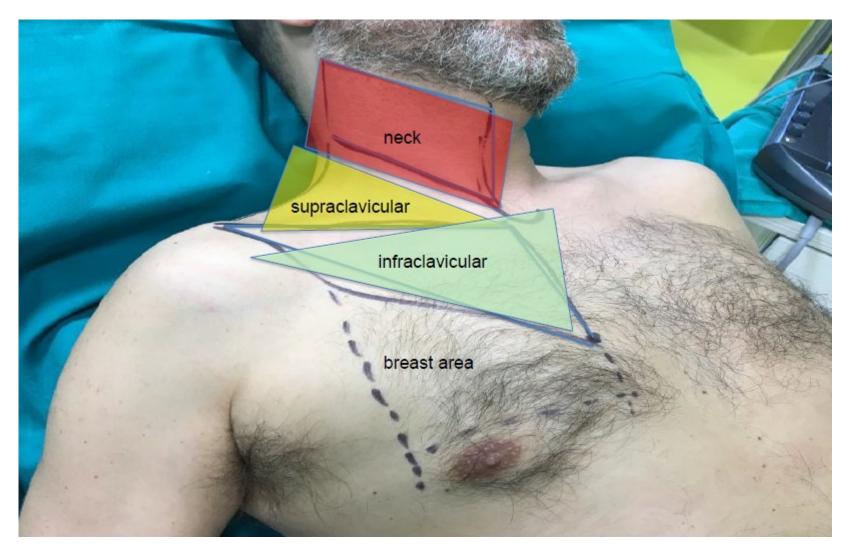
Figure 1. This person has a 21cm Total Zone Measurement (TZM), it divides into three 7cm zones to form the Red, Green and Yellow Zones. The ideal basilic vein image was located at 12cm from the medical epicondyle (MEC), in the Ideal Zone. Image by author.

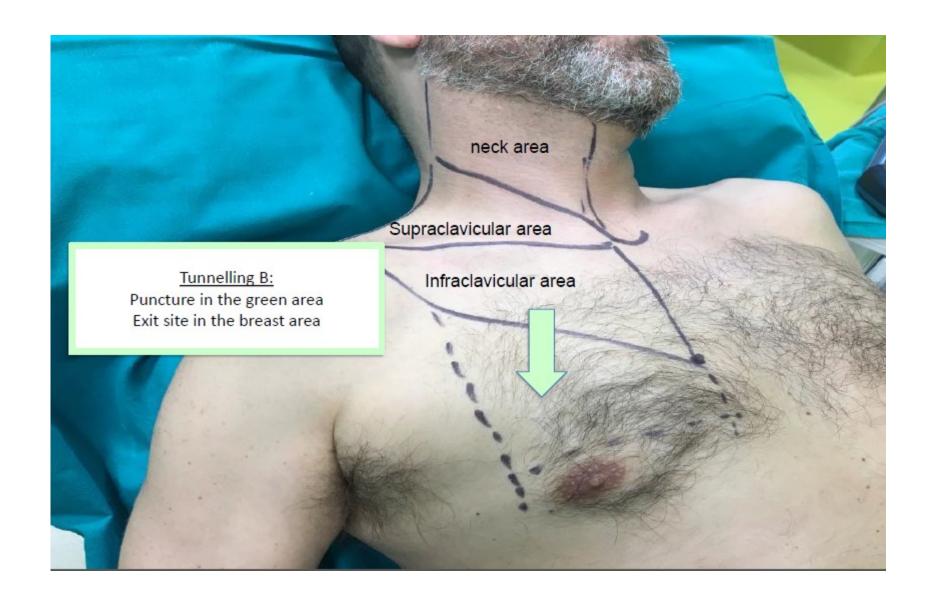


Bacterial skin colonization



ZIM for central venous access





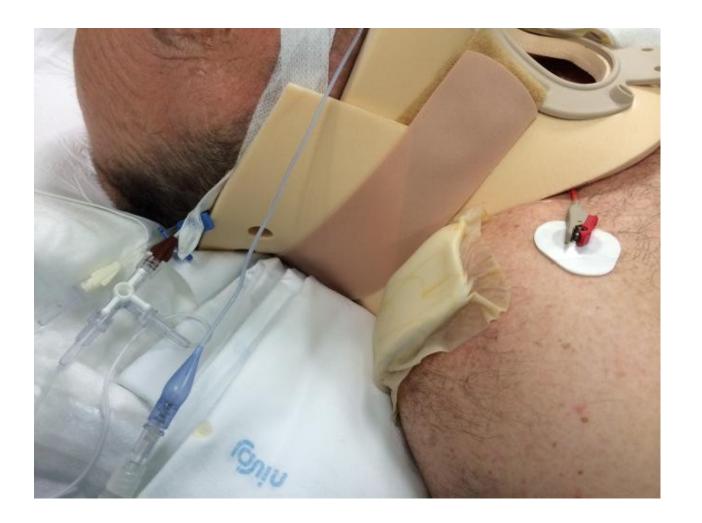
S laskavým svolením prim. Maňáska



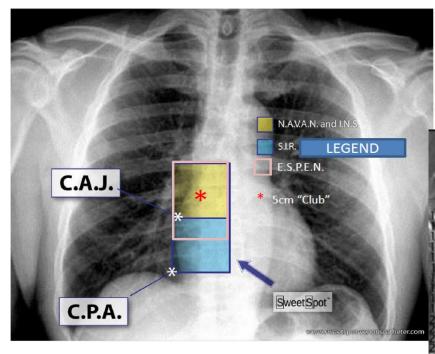
S laskavým svolením prim. Maňáska

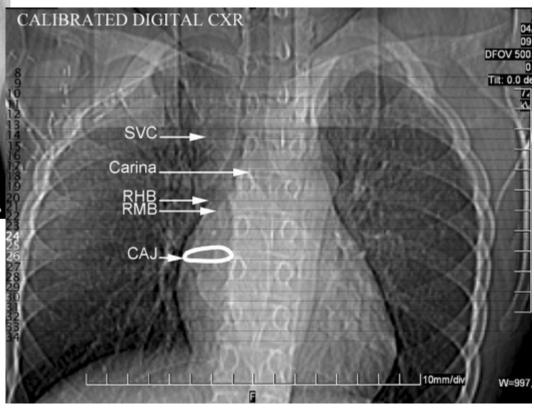


S laskavým svolením prim. Maňáska

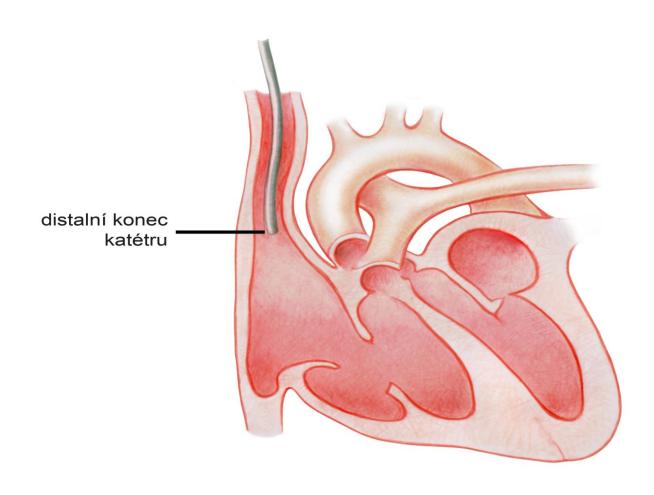


What "central" means?



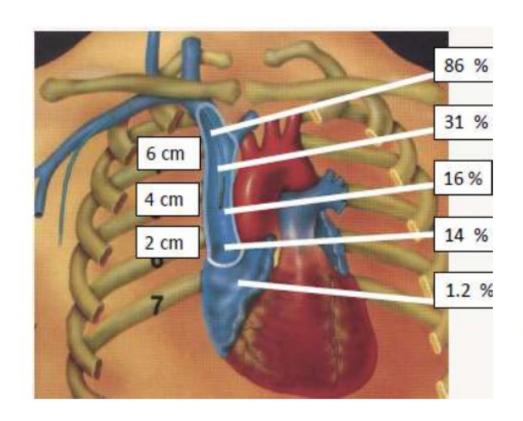


Distal tip of catheter



Silicone Venous Access Devices Positioned with Their Tips High in the Superior Vena Cava Are More Likely to Malfunction-Petersen





Durability proportional to depth

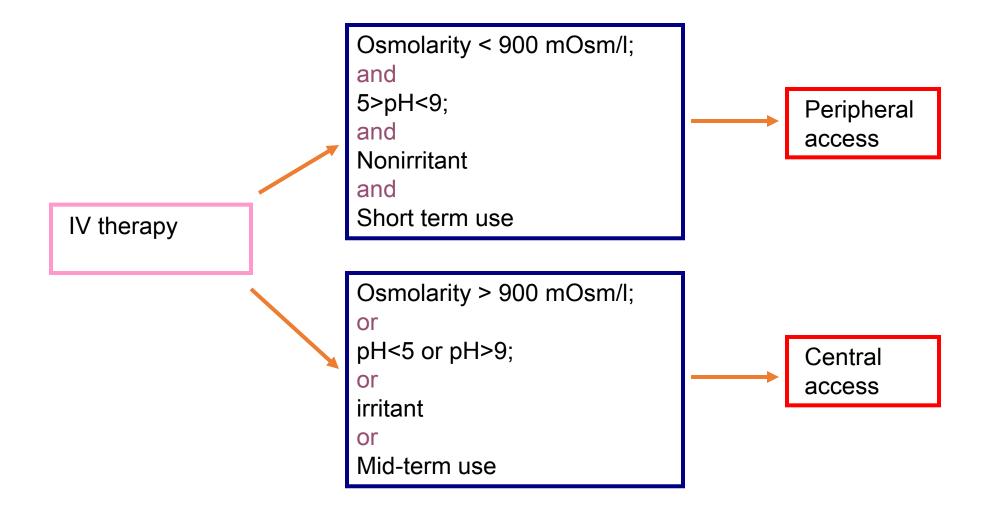
-RA 245d

-SVC/RA 116d

-SVC 100d

Source: Peterson et al. CVC Malfunction and Catheter's Tip Position. Am J Surgery 1999. Vol 178: 38-41

Peripheral or central access?



Complication when inserting long term access

Port, Broviac, PICC

- Failure to introduce
- Pneumothorax (not in PICC)
- Artery puncture
- Hematoma
- Nerve irritation
- Primary malposition
- Ductus thoracicus damaged (not in PICC)

Complication prevention

- Right indication of the type of venous access
- Appropriate vein and site of access (entrance and exit site)
- Ultrasound navigation
- Correct placement of tip of catheter

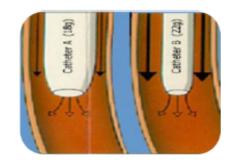
Xray check – necessary?

- US guidance is safe
- Experience of the performer
 -not necessary??
- Xray is a clear legal evidence of right position and absence of complications...

Catheter - vein ratio



PICC and vein ratio CVT Sharp R., 2015



Just enough to do the job!

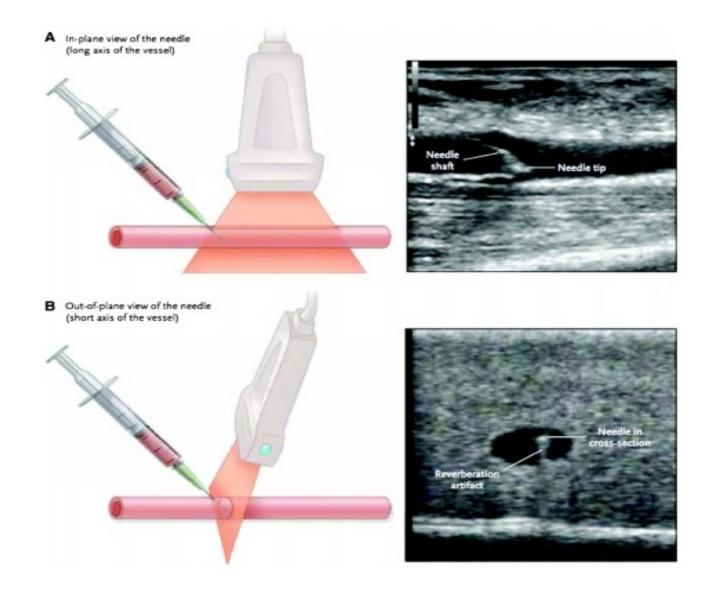
			4 Fr/1. 5mm diam.	5Fr/1 67mm diam.	6Fr/2r m diam.
	Diam.	Area	1 43mm ²	2.19mm ²	3.4mm ²
Vein	3.0mm	7.07mm ²	20%	31%	44%
	3.5mm	9.62mm ²	15%	23%	33%
	4.0mm	12.57mm ²	11%	17%	25%
	4.5mm	15.9mm ²	9%	14%	20%
	5.0mm	19.64mm²	7%	11%	16%
	5.5mm	23.76mm ²	6%	9%	13% 20

Catheter



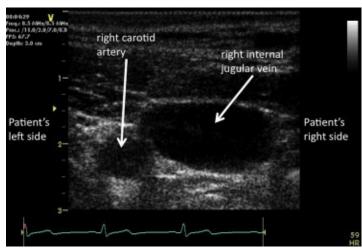
©Mari K. Cordes RN 5/2007

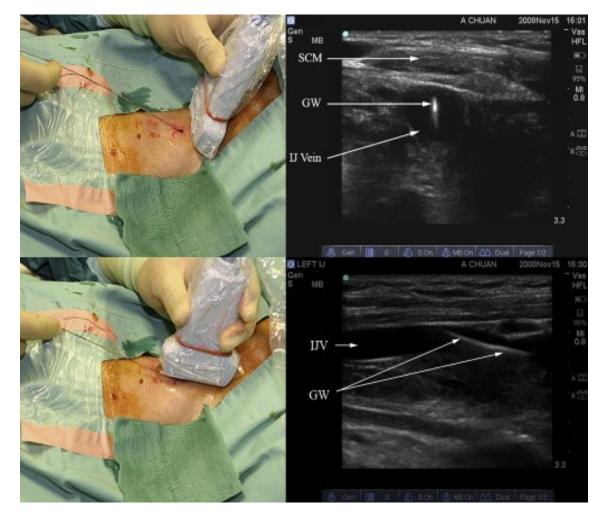
Ultrasound guidance

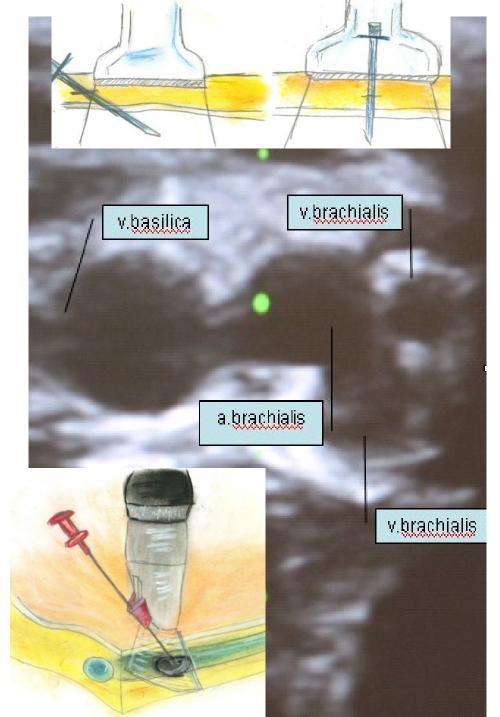


Ultrasound guidance













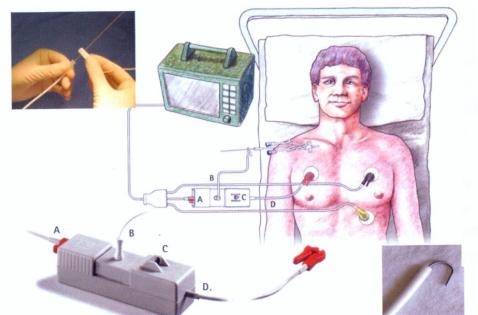
What is US good for?

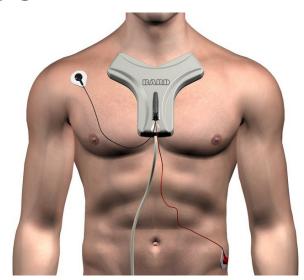
- Not for "hitting the right vein"
- rather to avoid complications and risks and malfuction immediately and in time

- choice of optimal site =
 - Better side? left/right
 - Best vein (lumen, unexpected anatomical structures, valves etc.)
 - Optimal entrance/exit site position
 - Check of the entire vein course...

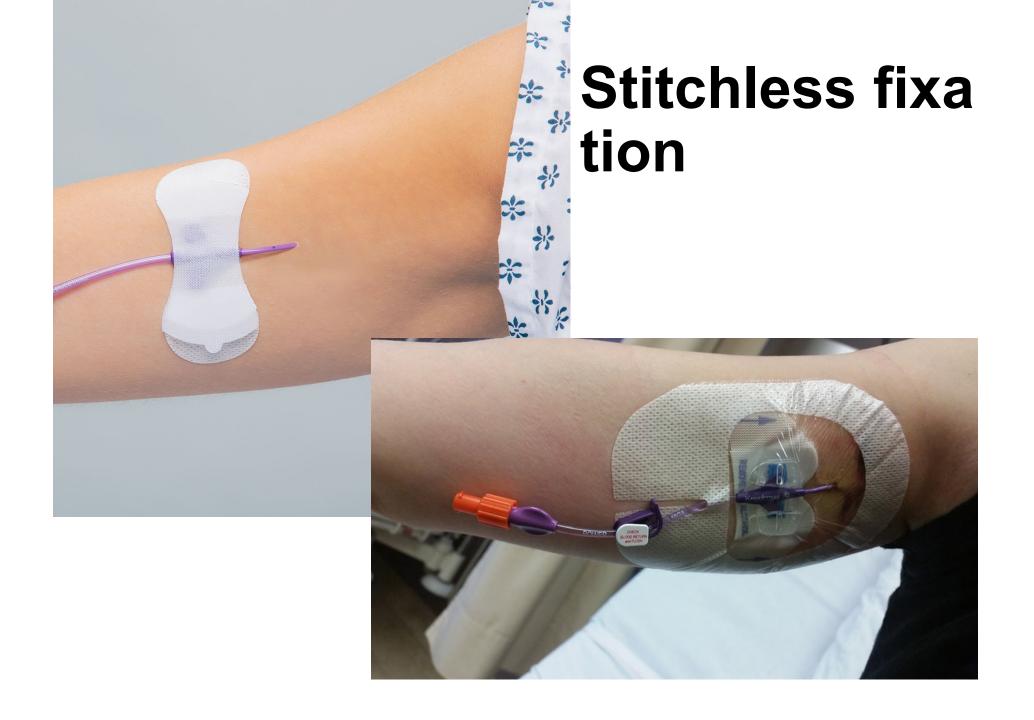
Tip position check?

- Estimate (measure on surface)
- Xray
- Intravasal ECG monitoring
- Magnetic guidance

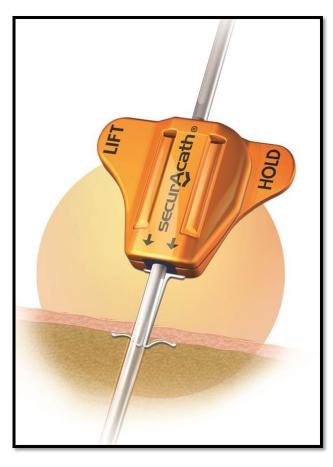


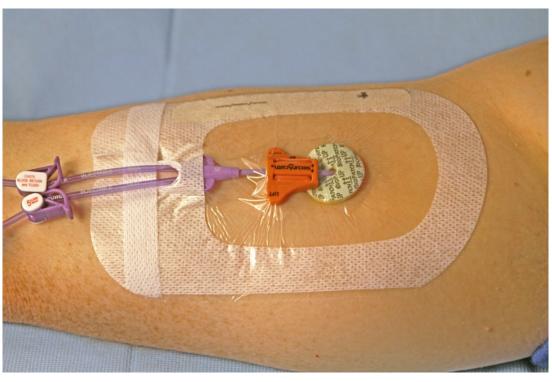


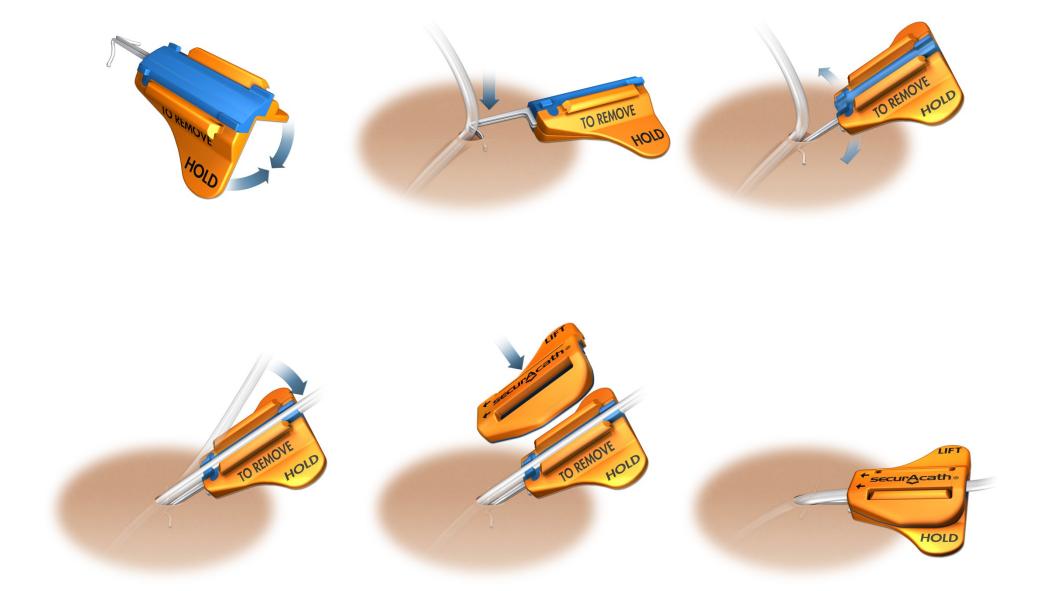




Stitchless fixation Securacath







Tissue glue (acrylic)



Complications -infection + thrombosis

- Peripheral cannula, CVC and midline
 - Always extract
 - (Pittiruti M, Hamilton H, Biffi R et al. ESPEN Guidelines on Parenteral Nutrition: central venous catheters (access, care, diagnosis and theraphy of complications) Clin Nutr 2009; 28:365-77)
- PICC
 - Local infection try to treat
 - Thrombosis- treat in situ, do not remove- full anticoagulation until the explantation
 - (Debourdeau P, Farge D, Beckers M et al. International clinical practice guidelines for the treatment and prophylaxis of thrombosis associated with central venous catheters in patients with cancer. J Thromb Haemost 2013; 11:71-80)
- Broviac, port
 - Try to treat and save

When to remove a Broviac or port

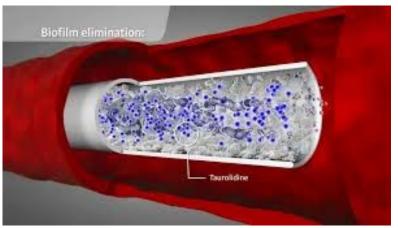
- Severe sepsis
- Tunnel infection (tunnelitis)
- Septic thrombosis
- Endocarditis
- Osteomyelitis
- Port chamber abscess
- Infection mycotic, Staph. aureus
 - G- bacteria, Staph koag. Neg. Or Enterococcus –treat
- AB i.v. for 2 weeks, endocarditis 6 weeks, osteomyelitis 8 weeks

Meemmel LA, Allon M, Bouza E et al. Clinical practice guidelines for the diagnosis and management of intravascular catheter-related infection: 2009 update by the Infectious Disease Society of America. Clin Infect Dis 2009; 49:1-45

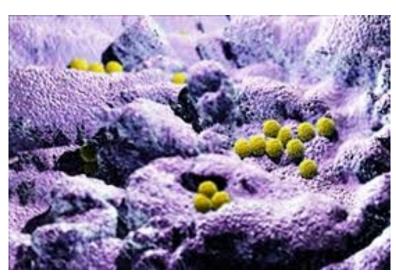
Fernandez-Hidalgo N, Almirante B, Calleja R et al. Antibiotic-lock theraphy for long-term intravascular catheter-related bacteremia: reset of an open, non-comparative study. J Antimicrob Chemother 2006; 57: 1172-80

Taurolidin in prevention of **biofilm** formation and its destruction









Complication prevention cont'd

Right indication

- Home or frequent administration- tunneled cath. Or port or picc
- Inpatients PICC better than untunneled catheter

Right flush technique

- start/stop method short bolus of saline repeated (2mls) producing turbulent flow
- Saline or taurolidin stopper (no more heparin)

PICC or port?

- indikation (who decides)?
- purpose
- Estimated time of use

- Nursing (PICC weekly, port every 6weeks)
- Swimming, sports, activities?
- Risks evaluation
- Availability (team, economics etc.)

PICC or port – length of use

- chemo 3-6 months (till a year)..... PICC
- chemo longer than 4-6 monthsport, PICC port

Economics (czk- ZUM)

PICC	port	PICC port
Cost 4884,-	Cost 5900,-	Cost 6400,-
(incl. Securacath, glue, ECG lead)	mikro 7700,-	
Implantation 1090 points	Approx. 1000 pts.	Approx. 1000 pts.

Nursing cost complex (material, time...)

PICC	port	
desinfekce, rukavice, tampony, proplach, lepení , bezjehlový konektor 53,50Kč	desinfekce, sterilní a obyč. rukavice, tampony, proplach, lepení, jehla 54,60Kč	
četnost 4x???	četnost 1x	
celkem 214,-Kč	celkem 54,60Kč	
	jehla s křidélky 117,50Kč	
vykazujeme ošetření PICC 0923745b.		
á týden???		
	odběr – jehla 30,-Kč	

Summary: what is right (state of the art) in venous access in 2023?

- Appropriate access, indication
- Right indroduction technique (US guided)
- Check of the catheter tip position
- Complication audit and multidisciplinary skilled and experienced team
- "vascular team"?



Right funcion, minimal burden

Thank you for attention

