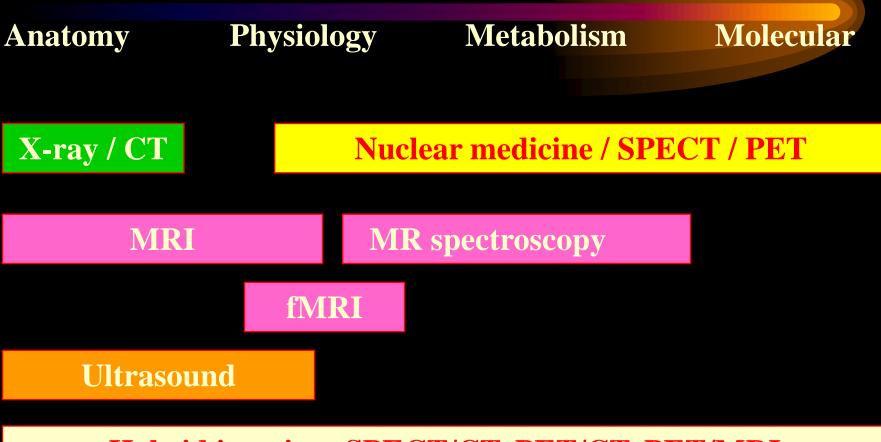
Nuclear medicine

Zámbó Katalin Department of Nuclear Medicine

Imaging tehniques



Hybrid imaging: SPECT/CT, PET/CT, PET/MRI

The short history of nuclear medicine

- Discovery of radioactivity (Bequerel 1896)
- Using of radioactive material as a tracer (György Hevesy 1923)
- Development of arteficial radioactivity (Irene Curie és Frederic Joliot Curie 1934)
- Gamma-camera (Anger 1951)



is the spontaneous desintegration (decay) of the nucleus of a radioactive atom, while the element becomes to an other one.

The hydrogen atom

THE BOHR MODEL OF THE HYDROGEN ATOM

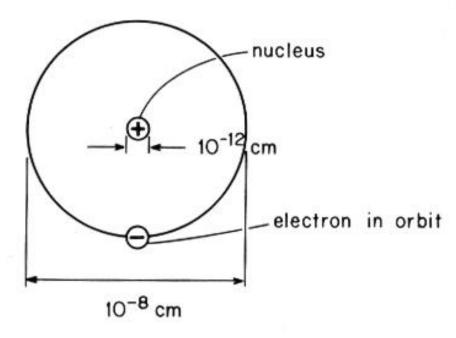


Fig. 1.1. The Bohr model of the hydrogen atom. The central nucleus contains essentially all the atom's mass, and is positively charged. The positive charge is balanced by the negative charge carried by the electron, which in this model circles the nucleus in a fixed orbit.

Sub-atomic particles

Table 1.1. PHYSICAL PROPERTIES OF SUB-ATOMIC PARTICLES

Particle	Electric Charge	Weight		Location
		Grams	a.m.u.	
Proton	+1	1.66×10^{-24}	1.0	Nucleus
Neutron	neutral	1.66×10^{-24}	1.0*	Nucleus
Electron	-1	9.1×10^{-28}	0.00054	Around nucleus

*The neutron is actually 0.08% heavier than the proton.

Number of protons = elemental identity number

Number of protons and neutrons = mass number

- Atoms with the same number of protons but differing number of neutrons are called isotopes of that element.

- The behaviour of the different radioactive isotopes of an element is the same as the stable form in every conditions.

Radioactive isotopes

Only certain combinations of protons and neutrons are stable, the other ones are radioactive, which become stable form with the emitting different radioactive radiations.

Activity

of a radioactive atom is usually given in desintegrations per second or minute, this is the *dps* or *dpm*.

The unit of the activity

- 1 Bq = 1 disintegration/second
- $1 \text{ kBq} = 10^3 \text{ disintegration/second}$
- 1 MBq = 10⁶ disintegration/second (used in practice)

Measurement

counts/second (cps) or counts/minute (cpm)

Half-life

is defined as the time required for one-half of the atoms in a group of radioactive atoms to decay.

- Physical half-life is characteristic for an element, independent on the external conditions.

- Biological half-life is depend on the physiological conditions (e.g. increased fluid input).
- Effective half-life: $1/T_{eff} = 1/T_{phys} + 1/T_{biol}$

Energy

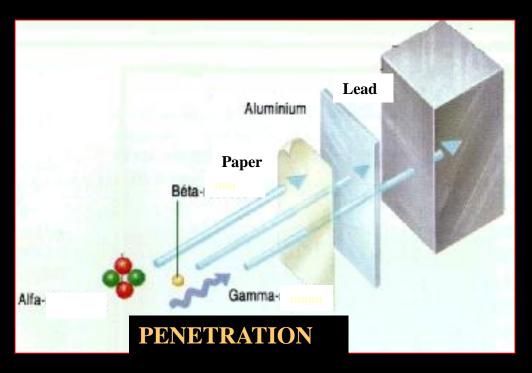
is emitted during the decay.

Units: eV, keV or MeV (1 eV is extremly small!)

Three kind of the radioactive radiation

- 1. Corpuscular :
- 2. Electromagnetic:

alpha -beta, +beta (positron) gamma



Alpha radiation

- the emission of a helium nucleus (2 protons + 2 neutrons) the ionizating property and biological effectivity is great - range in tissue is a few micrometers – can not be detected outside! -e.g. ²²⁶Radium for therapy (it is a new trend!)

Beta radiation

- the emission of high-speed electrons
- the biological effectivity is smaller than the alpha radiation
- the range in tissue is a few millimeters
- external detection is impossible, too
- the biological damage to tissues is high
- very suitable for radiotherapy
- e.g. ¹³¹Iodine for thyroid ablation

Gamma radiation

- really an electromagnetic radiation
- phisically similar to X-rays, but it comes from the nucleus of the atom
- very penetrated and easily pass trough tissue
- it can be detected externally well!
- e.g. 99mTechnetium for the diagnosis

The most commonly used isotopes

Isotope	Radiation	Half-time	Energy
99m-Technetium	γ	6 hours	140 kev
131-iodine	γ	8 days	364 keV
	(β		180 keV)
123-iodine	γ	13.2 hours	159 keV
111-indium	γ	2.8 days	172 keV
201-thallium	γ	3.1 days	76 keV

Equipments I.

Gamma-camera

- it "sees" the whole entire area below the detector



The layout of gamma-camera

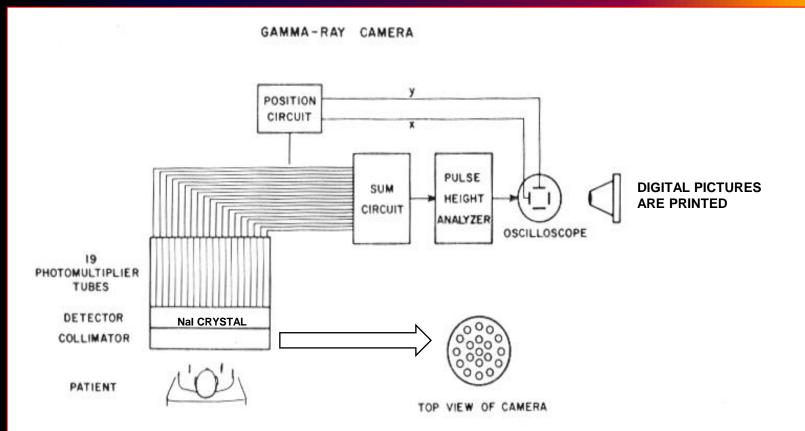


Fig. 1.11. The basic components of an Anger γ -ray camera. There is a one-to-one correspondence between the location of γ -ray interactions in the scintillation crystal and the location of the dot flashed on the oscilloscope screen.

Equipments II.

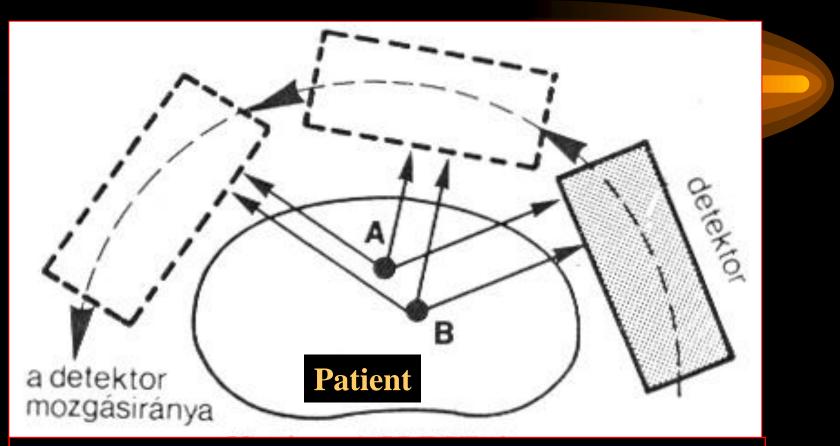
SPECT Single Photon Emission Computer Tomograph

SPECT/CT Multimodality!

- the computer program reconstruates the transversal, sagittal and coronal slices of the organ + fusion imaging



The principle of the SPECT



The detectors whirl around the patient and make pictures from different steps. The reconstruction and/or the reorientation are made by the computer program from this pictures after the imaging. Transversal, sagittal and coronal slices are reconstruated and evaluated.

+Beta (positron) radiation

– too many protons are in the nucleus

- its life is very short, when it slows down, it combines with a normal electron in a process known annihilation, which destroyes both electron and positron and produces two energetic gamma photons each with 511 keV

– isotopes with ultrashort half-life (11C, 15O, 13N, 18F) are used for PET examinations

– the metabolic changes of the tumors, the brain and the heart can be examined

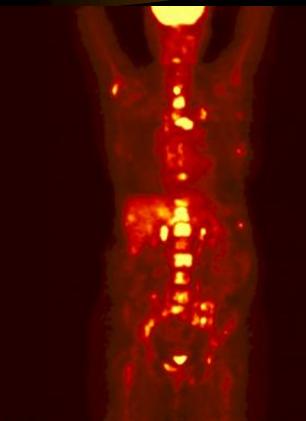
– e.g. 18Fluor-FDG shows the increased glucose metabolism of the tumors

Equipments III.

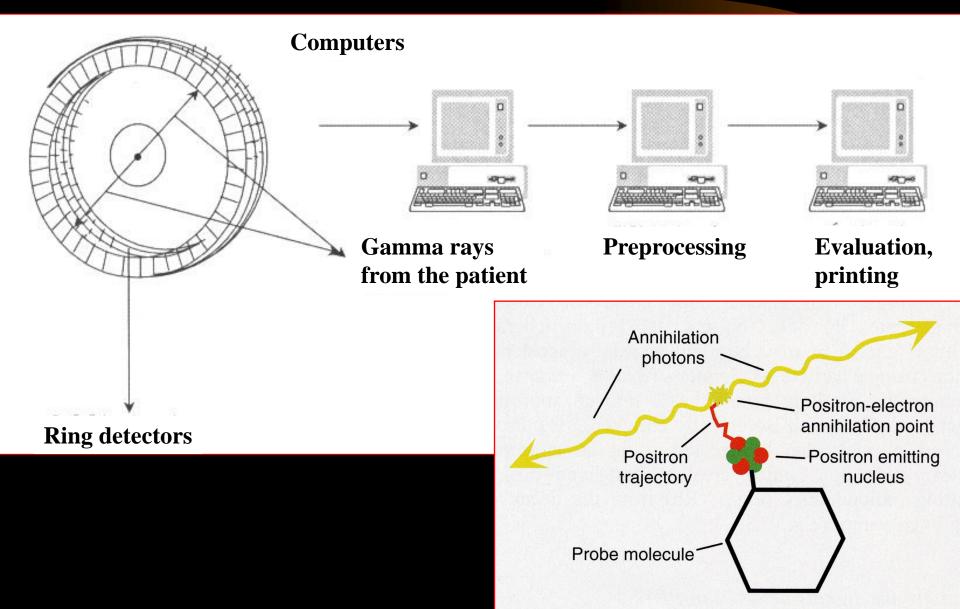
PET: Positron Emission Tomograph PET/CT: multimodality!



Multiplex bone metastases by FDG



The principle of the PET



Radiation exposure

principle of ALARA (as low as reasonable achieveble) both the patients and the staff
correct indication of the examination!
examination of pregnant women is contraindicated

- children should be examined carefully

Scintigraphies need

- gamma radiating isotope is detected by outside
- carrier molecule is participating in the examined function of the organs
- together is radiofarmaceutical
- administered in sterile intravenous NaCl injection
- delayed times are different before the examinations
- imaging by scintillation detector

In vivo radionuclide studies

- are based on the function of an organ or an organ system
- are very sensitive, but aspecific methods
- are easily performed
- need no premedication
- are not associated with any morbidity and complication, have only minimal risk
- are very good for screening studies

The types of the examinations

Static examinations (scintigraphy): - an optimal time-period after the subject administration is delayed and several photos are made of the organ from different directions

Dynamic studies:

- a frame-serie is stored in the computer from the time of the isotope injection during an optimal time-period of the examined organ function

Static examinations

- Thyroid with 99mTc-pertechnetate
- Lung with MAA (big particulums of HSA)
- **Bone** with MDP (methyl-diphosphonate)
- Bone marrow with Nanoalbumon (small particulated colloid)
- Liver and spleen with Fyton (big particulated colloid)
- Kidney with DMSA (dimercapto-succinate)
- Brain with DTPA (diethylen-triaminepentaacetate)

Dynamic studies

- Hepatobiliary scintigraphy
 Measurment of the hepatobiliary function from the blood through the liver to the bowels
- Camera-renography

Measurment of the renal function from the blood through the kidneys to the bladder

• Perfusion studies

Measurment of the perfusion of the several organs with fast excreted radiopharmaceuticals through the kidneys

Nuclear oncology

- Sentinel lymph node examination by human serum albumin (99mTc-Sentiscint)
- Neuroendocrine receptor study by 123I- or 131I-MIBG (pheochromocytoma, neuroblastoma)
- Somatostatin receptor study by 111Inoctreotide or 99mTc-depreotide (carcinoid tumors, small cell lung cancer, medullary thyroid cancer)

Sentinel lymph node examination

- Indications: mamma cancer
 - melanoma malignum
 - vulvar and penis malignancies
- Method: peritumoral injections by 4x15 MBq (4x0.2 ml) HSA colloid (99mTc-Sentiscint)

 static images from the lymph nodes

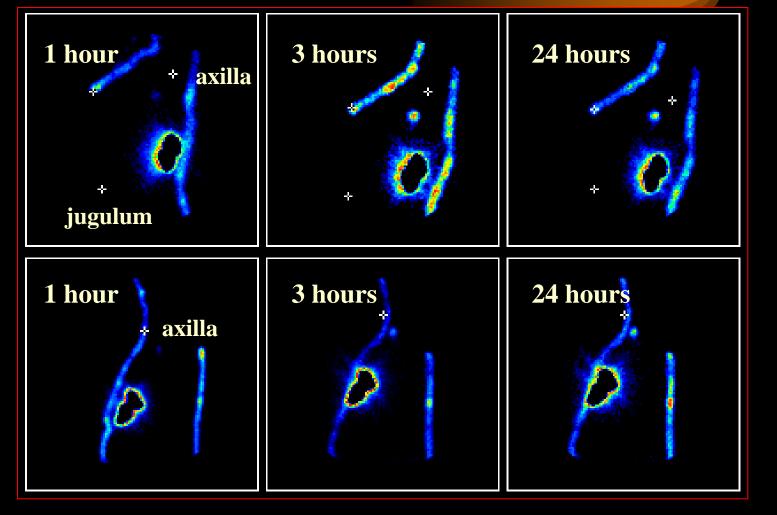
 1, 3 és 24 hours after the injection

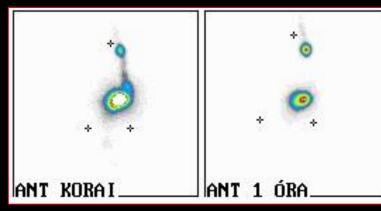
 the sentinel lymph nodes are marked
 on the skin, the operation is on the following
 day with help of *intraoperative gamma-probe*

Sentinel lymph node scintigraphy in breast cancer in the left side

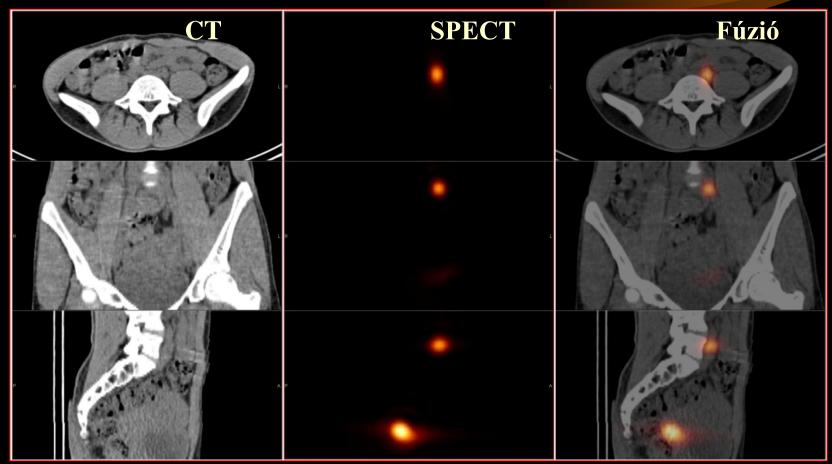
Anterior







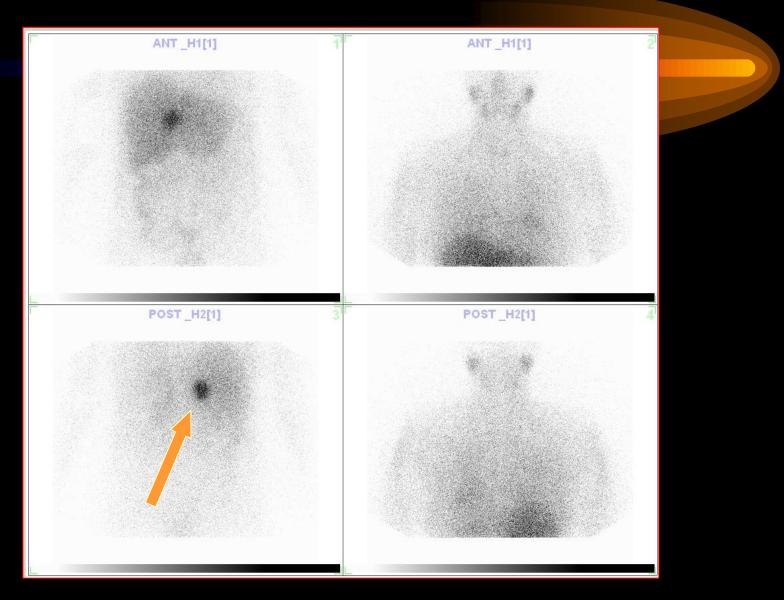
Sentinel lymph node scintigraphy in adenocc cervicis uteri in the left parailiacalis region



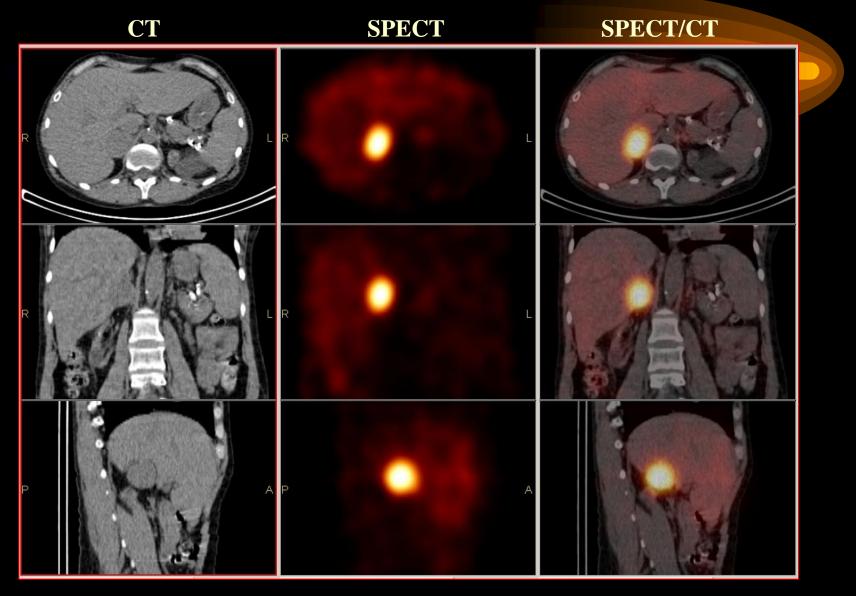
Adrenerg receptor scintigraphy

- Injected subject: 185 MBq 123-iodine-MIBG (metaiodobenzyl-guanidine) is binding to adrenerg receptors of the tumor-cells
- **Imaging time:** 6 és 24 hours after the intravenous injection (SPECT/CT imaging!)
- Indications: neuroendocrine tumors
 - pheochromocytoma
 - neuroblastoma

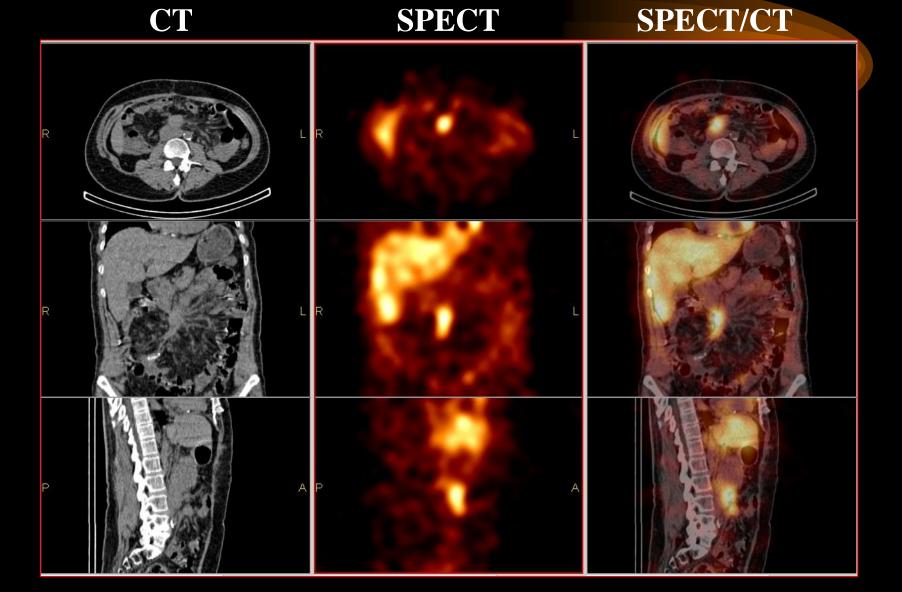
Phaeochromocytoma in the right adrenal gland



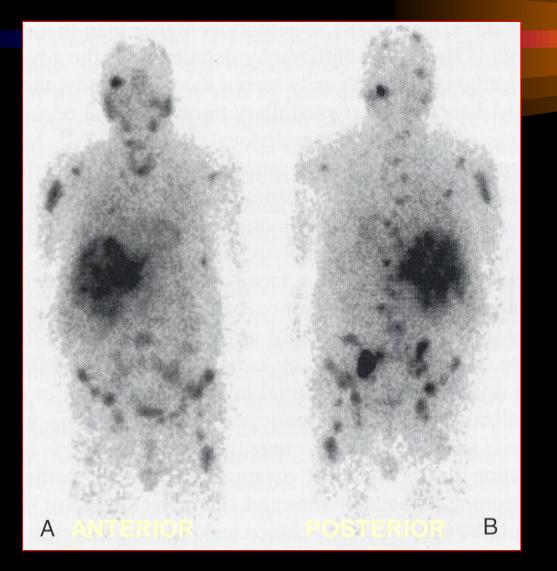
Phaeochromocytoma in the right adrenal gland by SPECT/CT



Metastases in retroperitoneal lymph nodes after operation of small intestine NET



Multiplex 123-iodine-MIBG cumulation in malignant pheochromocytoma



Somatostatin receptor scintigraphy

• Injected subject: 122 MBq 111-Indium-pentetreotide or 740 MBq 99m-Tc-depreotide

(somatostatin analog peptids are binding to the receptors overexpressed on the surface of tumor cells)

Imaging time: - 99m-Tc on the same day 2 hours later
 - 111-In 24 and 48 hours

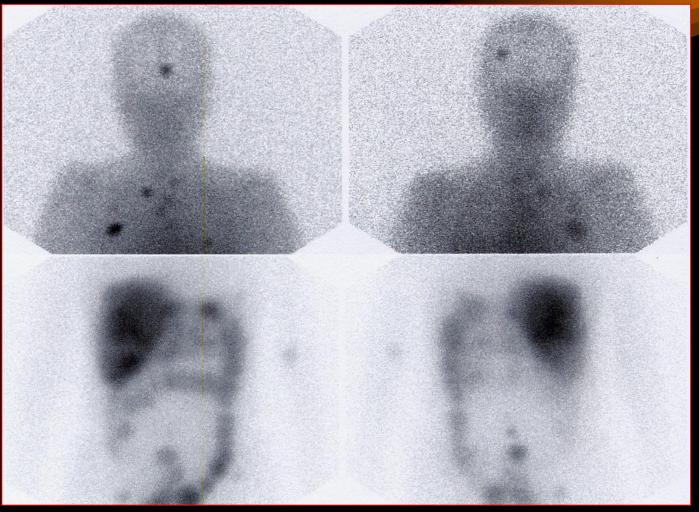
after the intravenous injection (SPECT/CT imaging!)

- Indications: carcinoid és (
 - carcinoid és GEP tumors
 - small cell lung cancer
 - medullary thyroid cancer

St. p. pancreas head carcinoid operation, metastases? 111In-Octreoscan-study

ANTERIOR



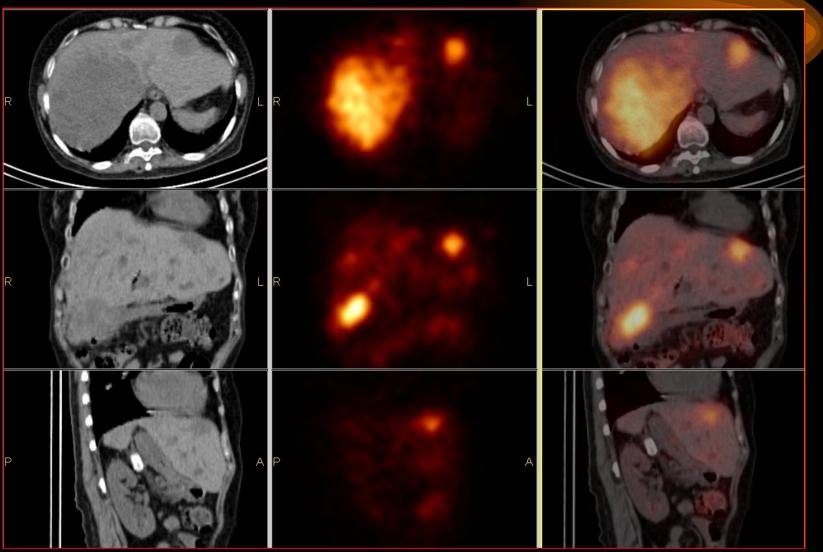


Multiplex liver metastases of carcinoid

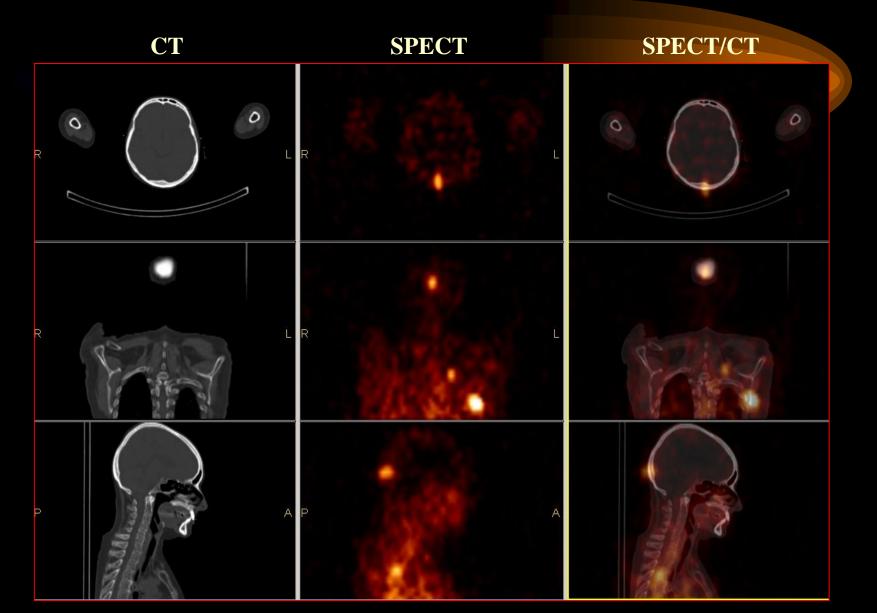
CT

SPECT

SPECT/CT

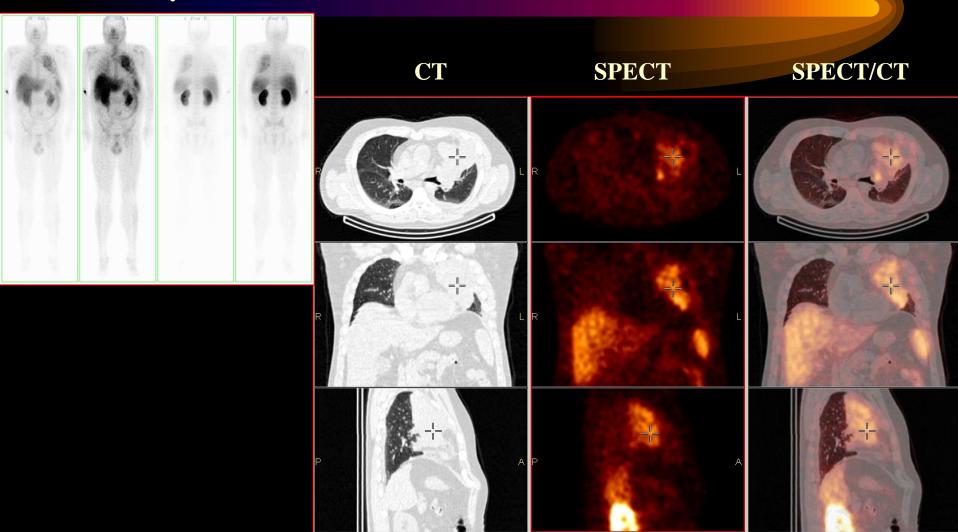


Multiplex bone metastases of carcinoid

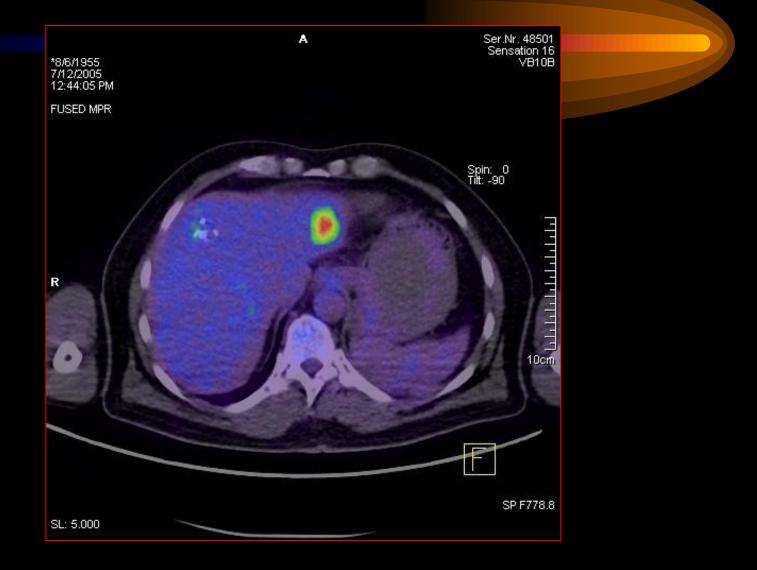


Carcinoid in the left lung? 99mTc-Neospect examination

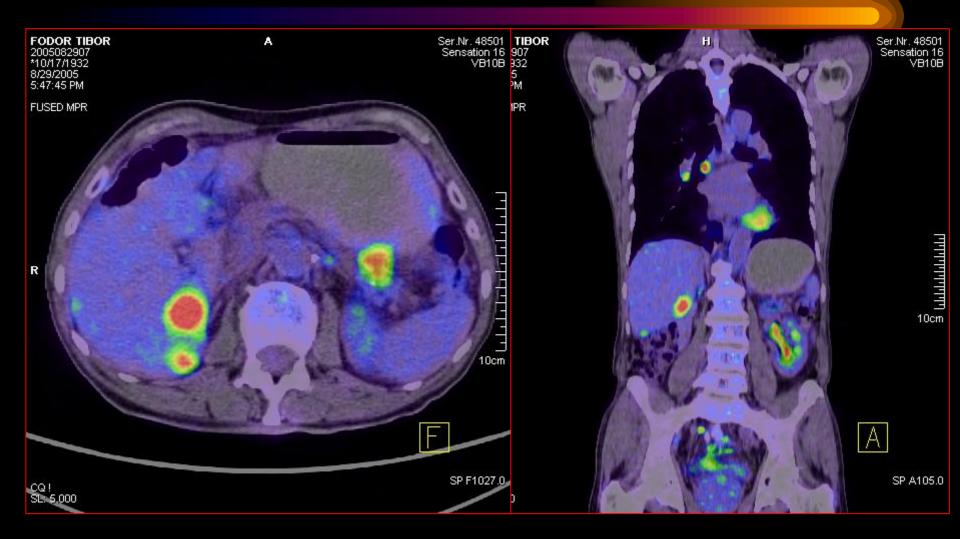
Whole body scan



Liver metastasis of rectal cancer by 18F-FDG



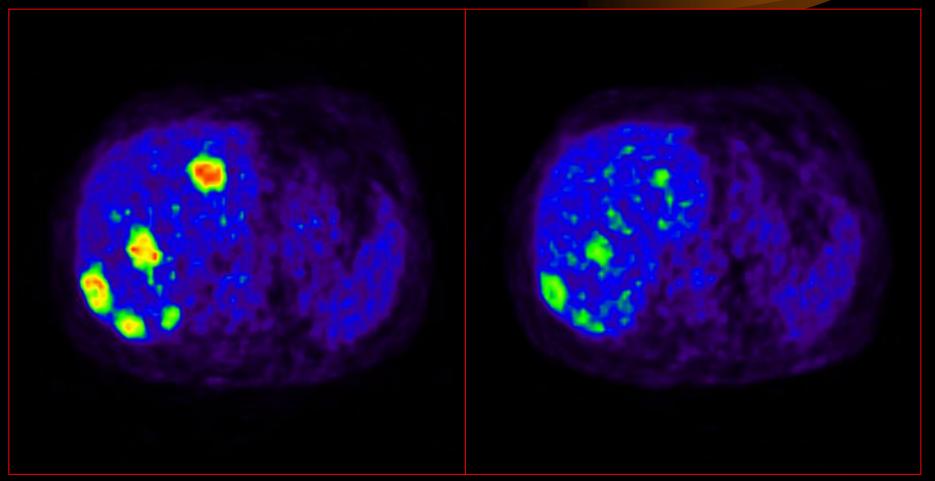
Multiplex metastases of pancreas tail cancer by 18F-FDG



Multiplex liver metastases of sigmatumor by 18F-FDG

Before therapy

After therapy



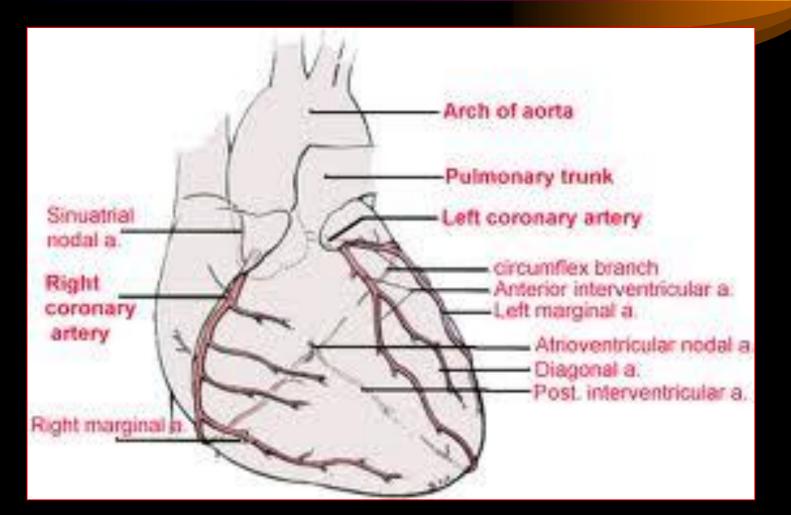
Nuclear cardiology

- Rest myocardial perfusion study
- Stress/rest myocardial perfusion study
- Radionuclide ventriculography (RNV), multigated analysis (MUGA)

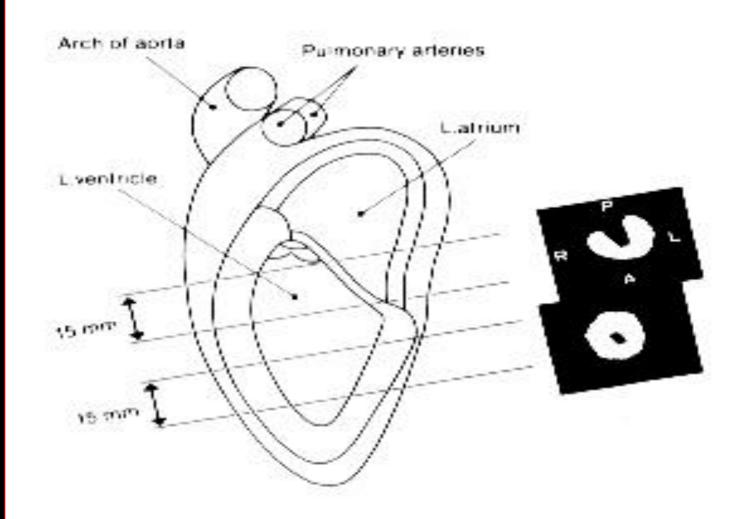
Myocardial perfusion imaging in rest conditions

- The myocardium is shown by radioactive tracers (99mTc-MIBI, 99mTc-tetrofosmin, 201-Tl-clorid)
- Reconstruated and reorientated slices are investigated from the left ventricle by SPECT
- The impairment of the myocardial perfusion is indicated by decreased activity or lack of the activity

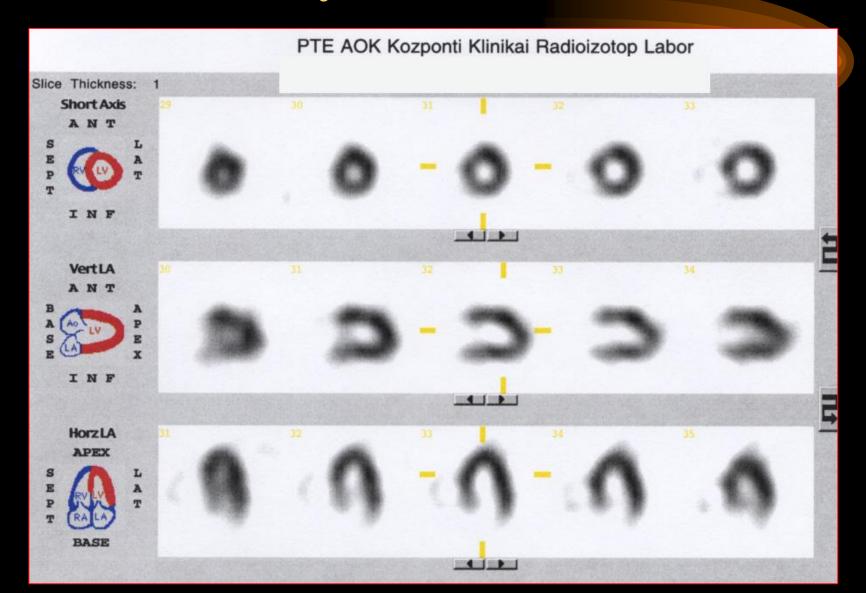
Coronary anatomy



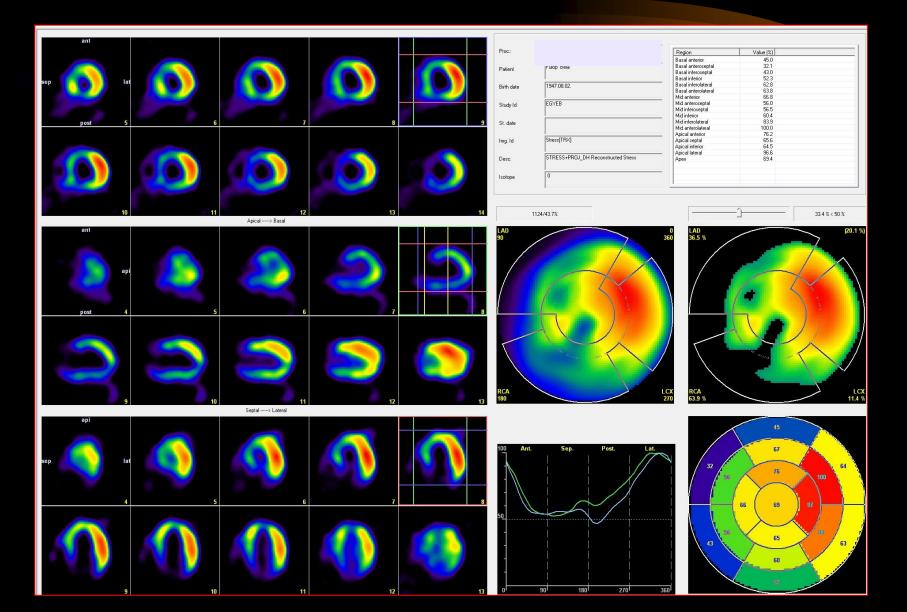
Long axis and short axis slices of the myocardium by SPECT



The transversal, sagittal and coronal slices of the myocardium



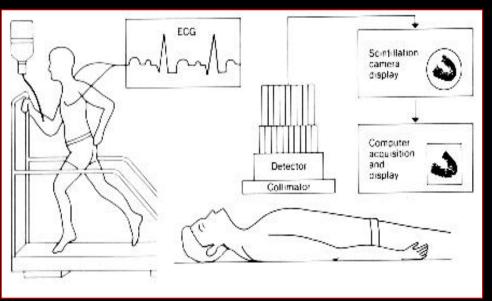
Infero-septal + antero-septal hypoperfusion



Stress/rest myocardial perfusion study

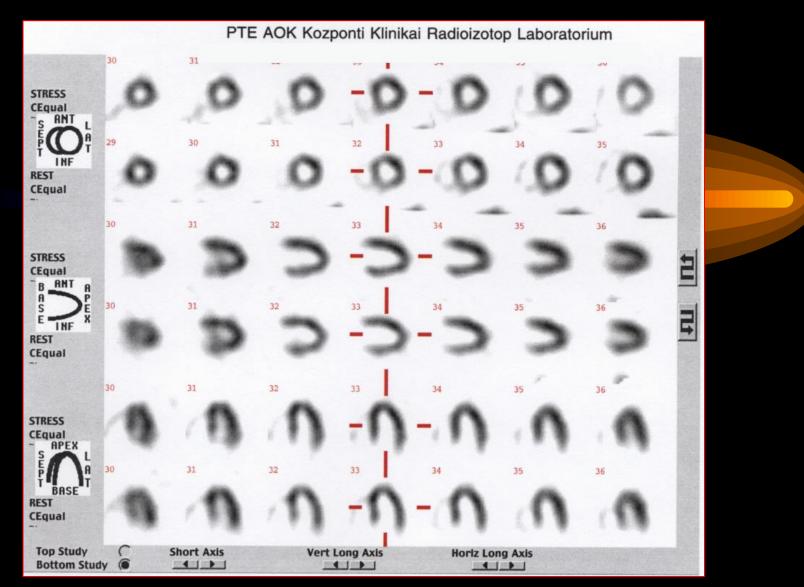
- Physical or pharmacological stress (Dipyridamol, Dobutrex) is applied
- The isotope is administered on the top of the stress » *SPECT-imaging*
- Rest SPECT-imaging is on the same day (Tl), or a day later (Tc-MIBI)
- Evaluation by two independent nuclear medicine experts + cardiologist

The method of the imaging by fixed 90 degree double-haed SPECT (Physical or Dipyridamol stress is used commonly)

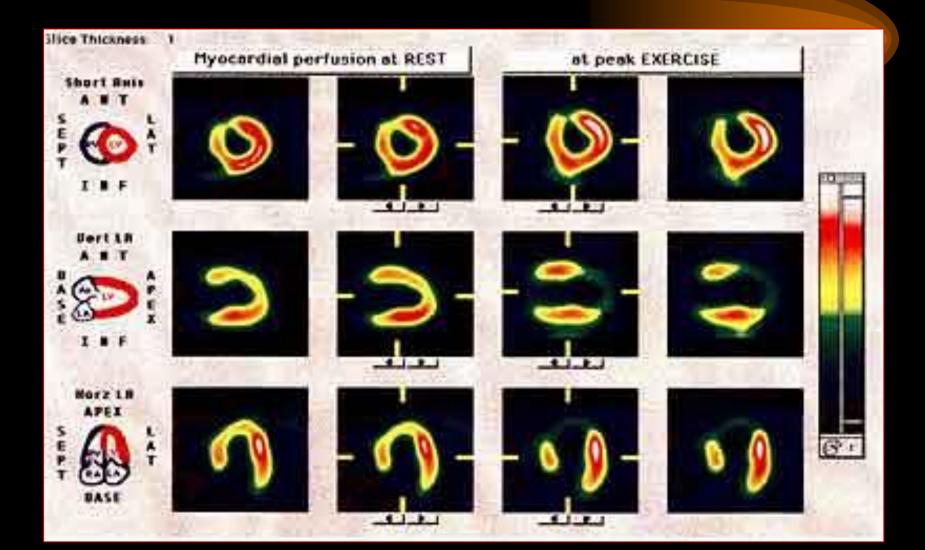




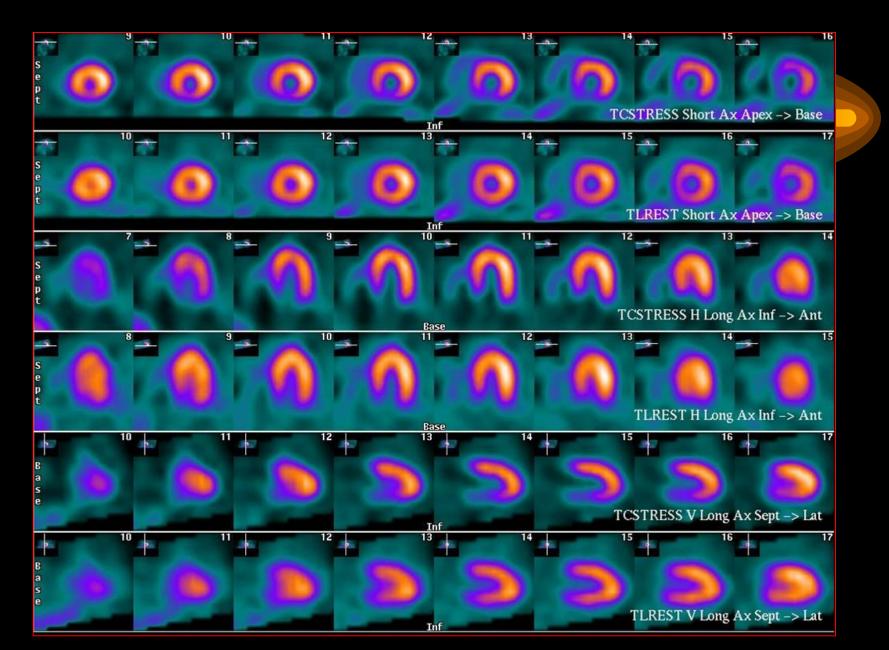
Normal stress/rest myocardial perfusion study



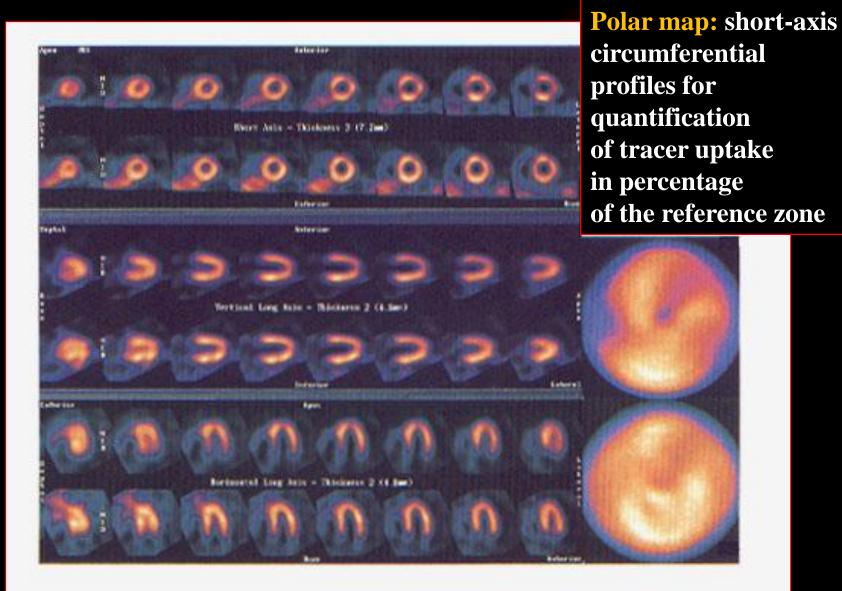
Transient ischaemy in the apex and in the apical part of the antero-septal wall



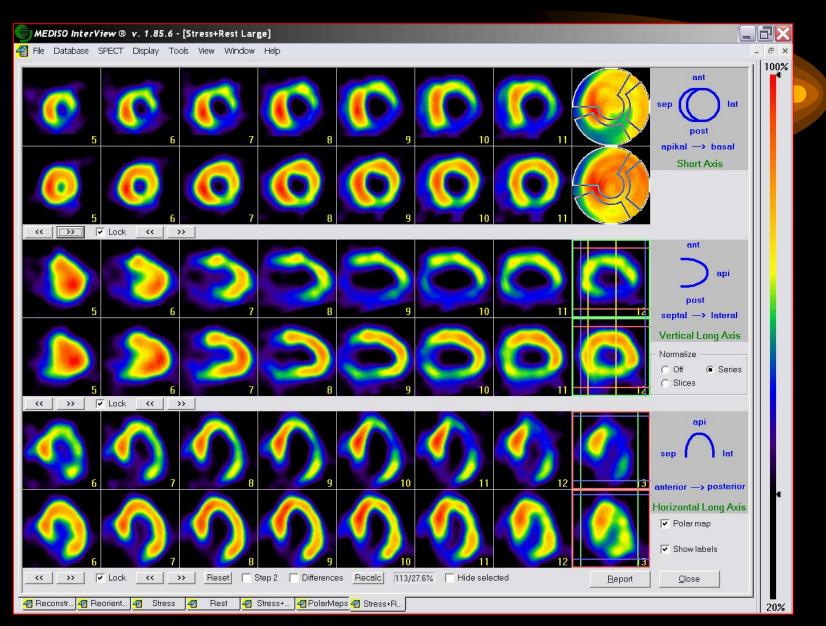
CAD in the infero-septal wall



CAD in the basal part of the septum



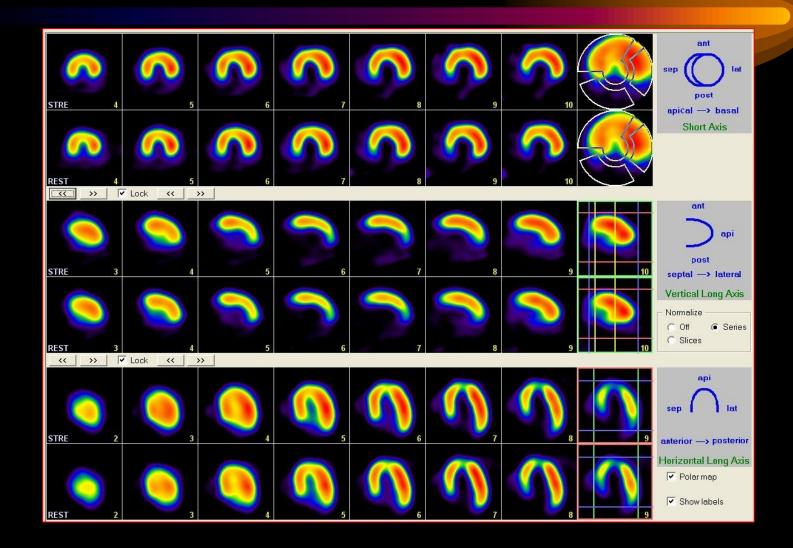
Severe infero-lateral transient ischaemy



Severe infero-lateral transient ischaemy – 3D processing

🗄 PerfSPECTive(TM) Display - The Emory Cardiac Toolbo	ox(TM) - Copyright 1998			
File Options Gated PerfSPECTive(TM) Parameters Slices Polar Maps	Match/Mismatch Tool Perf	SPECTive(TM) Functional Analysis	Summary Page Save & Exit	
Back of the second seco	Blackout	Reverse	Administrate Save a Lit	

Scar in the inferior, infero-septal and infero-lateral wall



Viability of the myocardium

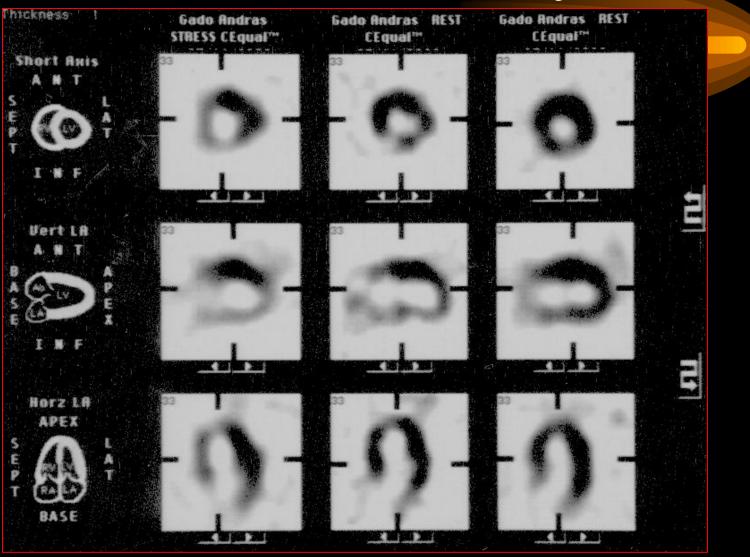
• When fix defect is found in both stress and rest situation (scar or hibernated myocardium) to assess the possibility for succesful revascularization

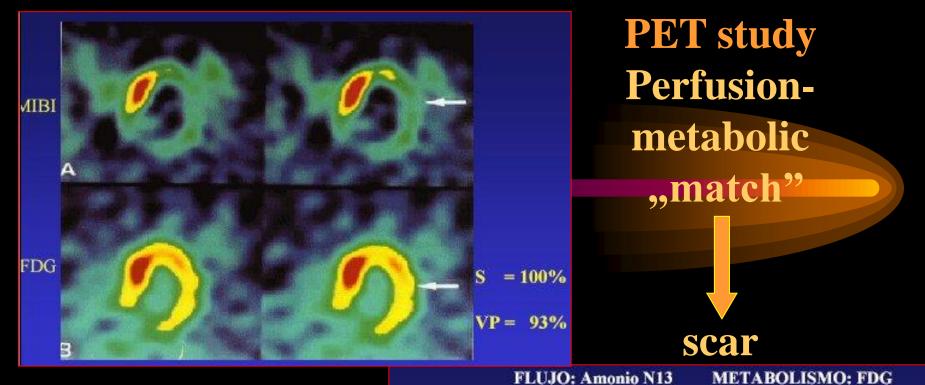
• 201Tl-chlorid has a specific redistribution pattern after 3-4 hours in rest, which depends on the wash-out from the myocytes

• After the reinjection the activity of the myocardium depends on primarily the perfusion by the coronary arteries

Viability examination by 201Tl-clorid

stress redistr. reinjection



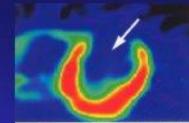


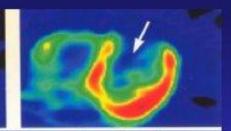
MIBI-FDG "mismatch"

myocardium

is viable

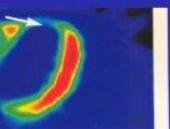
Match concordancia





No viable territorio Arteria Descendente Anterior

Mismatch discordancia



0

Viable territorio Arteria Descendente Anterior

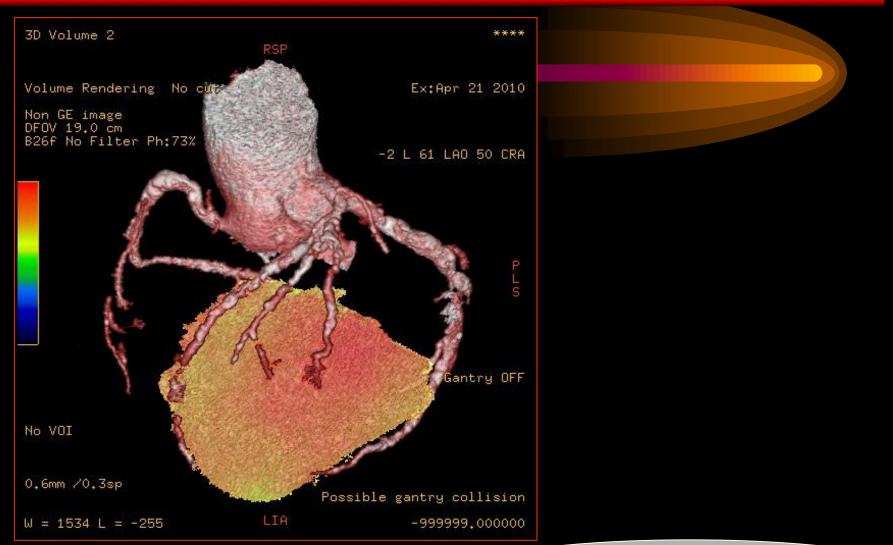
3D SPECT/CT imaging:

stenosis of right coronary artery

apical hypoperfusion







Balogh I. – Kerecsen G.

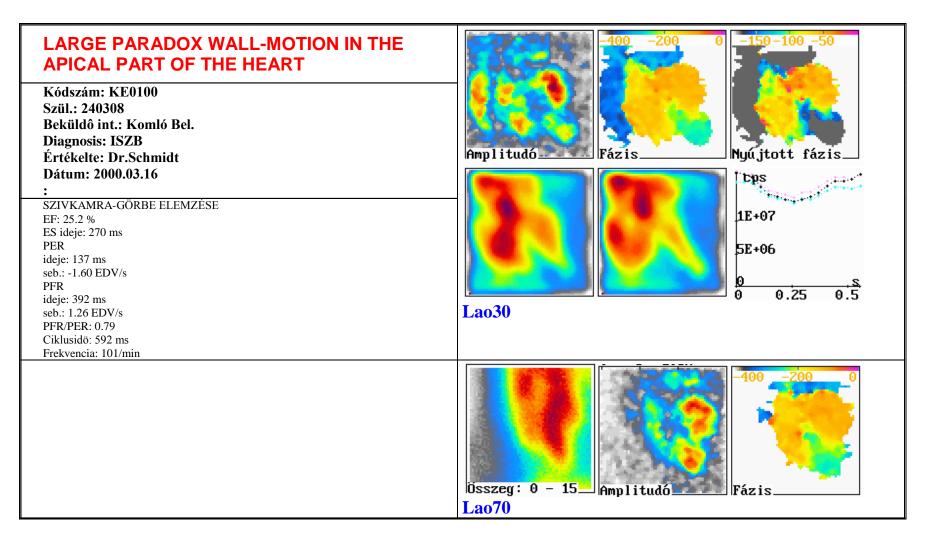
Radionuclide ventriculography (RNV), multigated analysis (MUGA)

- The blood-pool of the heart is labelled (99mTc-pyrophosphate-RBC)
- Gamma-camera-computer-R wave monitor system
- **EF=ED ES/ED-BG (%)**
- Wall-motion is analysed by parametric pictures

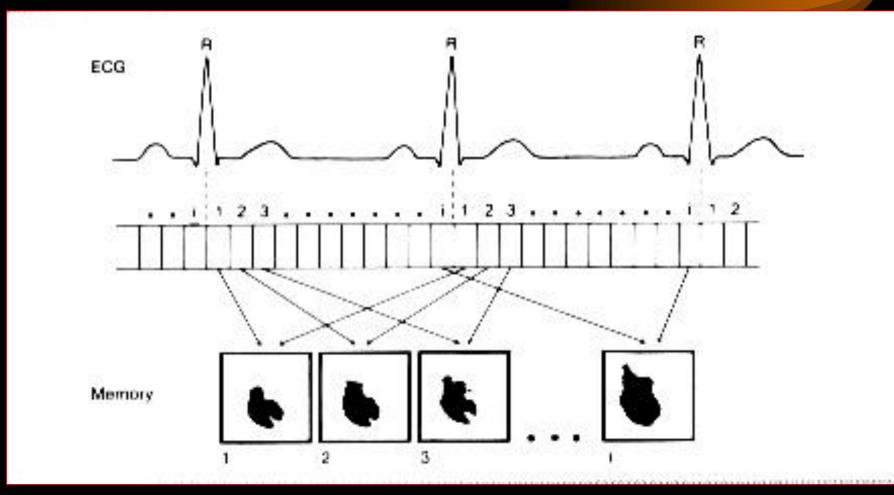


VENTRICULO SZCINTIGRÁFIA EREDMÉNYLAP

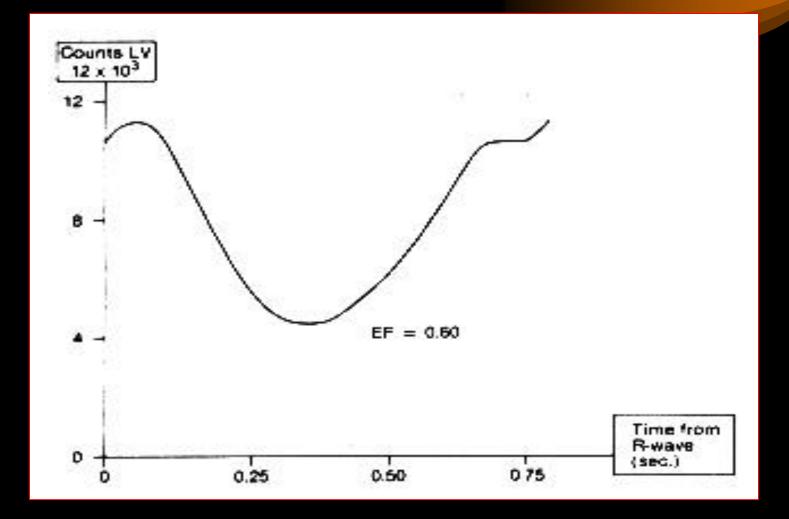
PÉCSI TUDOMÁNYEGYETEM ÁLTALÁNOS ORVOSTUDOMÁNYI KAR Központi Klinikai Radioizotóp Laboratórium 7624 Pécs, Ifjúság útja 13. Tel.: (72) 326-222/1229 Intézetvezető: dr. Zámbó Katalin



An average cycle is generated from more hundred heart cycles



The ejection fraction curve





VENTRICULO SZCINTIGRÁFIA EREDMÉNYLAP

PÉCSI TUDOMÁNYEGYETEM ÁLTALÁNOS ORVOSTUDOMÁNYI KAR Központi Klinikai Radioizotóp Laboratórium 7624 Pécs, Ifjúság útja 13. Tel.: (72) 326-222/1229 Intézetvezető: dr. Zámbó Katalin

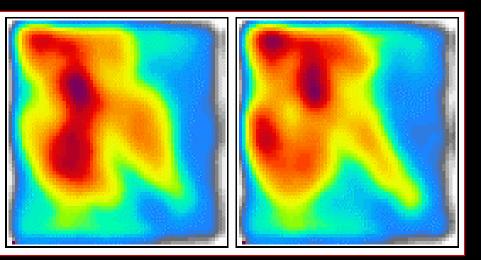
NORMAL FUNCTION OF THE LEFT VENTRICLE Kódszám: KE0351 Szül.: 450515 Beküldô int.: Szigetvár Bel. Diagnosis: St.p.inf.myoc. Értékelte: Dr.Schmidt Dátum: 2000.10.02	$ \begin{array}{ c c } \hline \hline \\ $
: SZIVKAMRA-GÖRBE ELEMZÉSE EF: 64.1 % ES ideje: 398 ms PER ideje: 180 ms seb.: -2.50 EDV/s PFR ideje: 550 ms seb.: 2.18 EDV/s PFR/PER: 0.87 Ciklusidö: 944 ms Frekvencia: 64/min Infl. pont: 768 ms	Végdiasztole Vágdiasztole
	$\begin{bmatrix} 1 & 1 & 1 \\ 0 & 1 & 0 \\ 0 & 1 & 0 \end{bmatrix} \begin{bmatrix} 1 & 1 & 1 \\ 0 & 1 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 1 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 &$

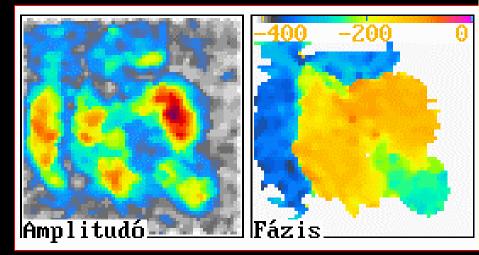
Parametric pictures

Amplitude picture

The colours represent the amplitude of the change of the activity of the pixels. Phase picture

The colours represent the phase of the change of the activity of the pixels.







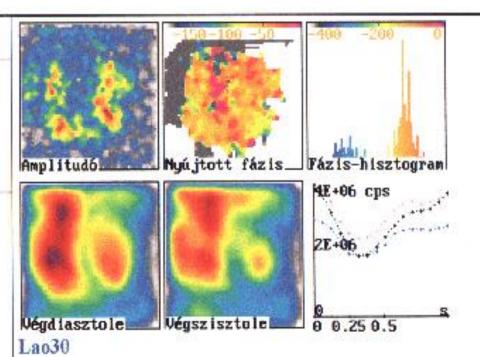
VENTRICULO SZCINTIGRÁFIA EREDMÉNYLAP

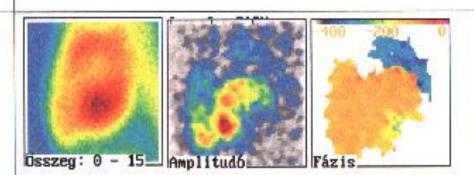
PÉCSI TUDOMÁNYEGYETEM ÁLTALÁNOS ORVOSTUDOMÁNYI KAR Központi Klinikai Radioizotóp Laboratórium 7624 Pécs, Ifjúság útja 13. Tel.: (72) 326-222/1229 Intézetvezető: dr. Zámbó Katalin

POSTERO-INFERO-LATERALIS HYPOKINESIS

Kódszám: KE0082 Szül.: 50.12.08. Beküldő int.: Szigetvár Kard.Szakr. Diagnosis: St.p.AMI Értékelte: dr.Udvaros Dátum: 2000.03.01

SZIVKAMRA-GÖRBU ULUMZÉSE EF: 52.6 % ES ideje: 378 ms PUR ideje: 190 ms seh: -2.53 hDV/s PUR ideje: 544 ms sen: 1.34 EDV/s PUR/PUR: 0.53 Ciklusidő: 1024 ms Frekvencia: 59/min Infl. pont: 803 ms







VENTRICULO SZCINTIGRÁFIA EREDMÉNYLAP

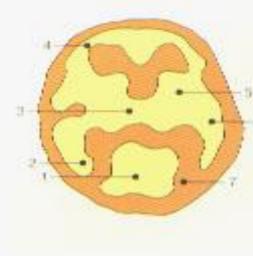
PÉCSI TUDOMÁNYEGYETEM ÁLTALÁNOS ORVOSTUDOMÁNYI KAR Központi Klinikai Radioizotóp Laboratórium 7624 Pécs, Ifjúság útja 13. Tel.: (72) 326-222/1229 Intézetvezető: dr. Zámbó Katalin

LARGE HYPOKINESIS IN DCM	
Kódszám: KE0156	
Szül: 330801	
Beküldő int.: PTE II.Bel.kl. Diagnosis: DCM	
Értékelte: dr.Schmidt	
Dátum: 2000.04.19	Amplitudó Fázis Nyújtott fázis
SZIVKAMRA-GÖRBE ELEMZÉSE	
EF: 28.5 % ES ideje: 346 ms	ZE+07
PER	
ideje: 194 ms	
seb.: -1.30 EDV/s PFR	March 1997 A Street Stree
ideje: 433 ms	
seb.: 0.35 EDV/s PFR/PER: 0.27	Végdiasztole Végszisztole 0 0.25 0.5
Ciklusidö: 832 ms	Lao30
Frekvencia: 72/min	
Infl. pant: 659 ms	
	Osszeg: 0 - 15
	Lao70

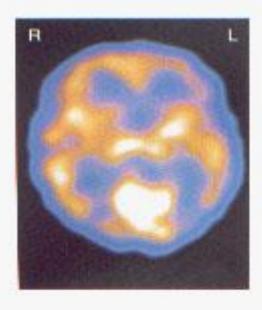
Brain perfusion study

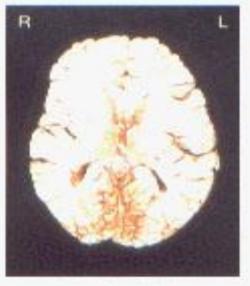
- The cortex and the basal ganglia are shown by a lypophil radioactive subject (99mTc-HM-PAO – hexamethyl-propilenamine-oxyme)
- Reconstruated and reorientated transversal, sagittal and coronal slices from the brain
- The impairment of the brain perfusion is indicated by decreased activity or lack of the activity

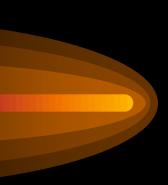
The parts of the brain



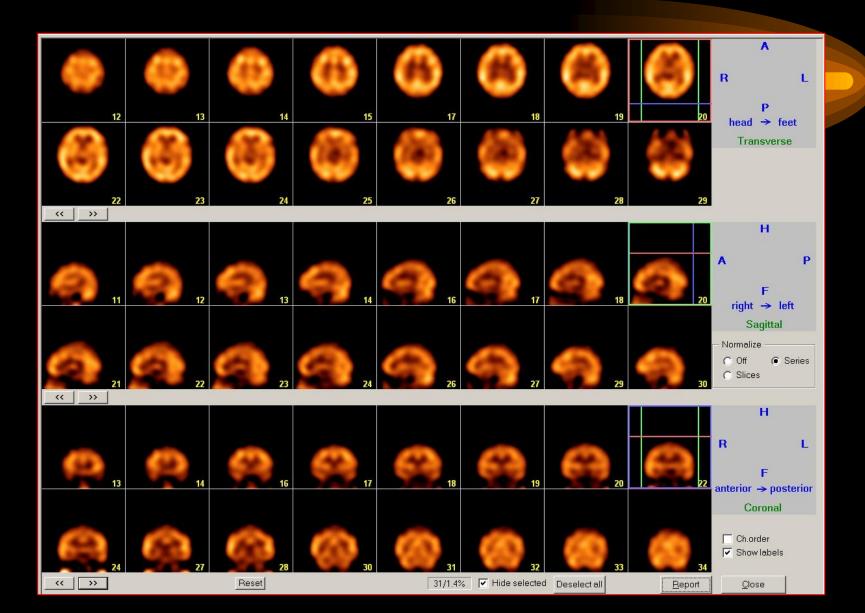
Seeing cortex
 Occipital lobe
 Thalamus
 Frontal lobe
 N. caudatus
 Temporal lobe
 Chambers

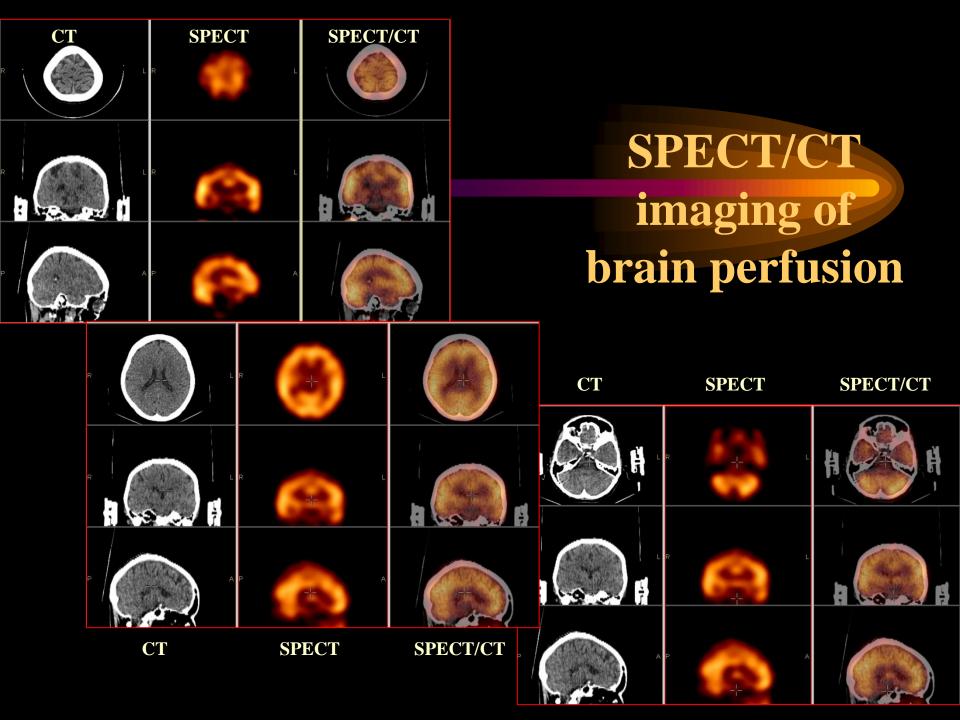




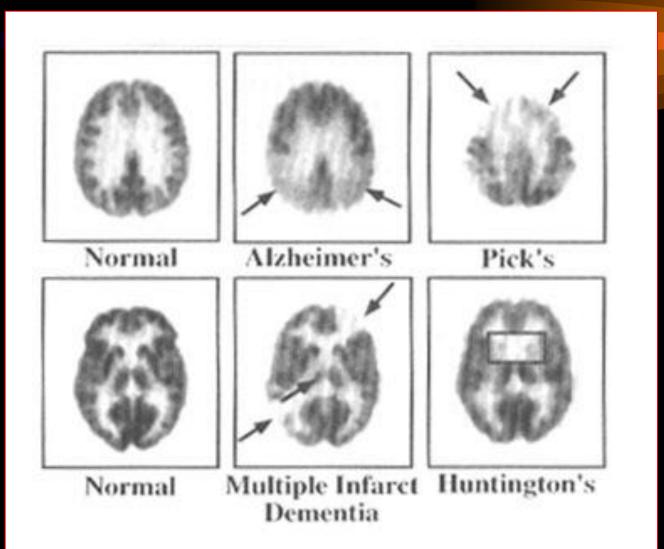


Normal brain perfusion

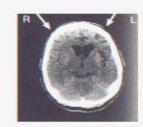




The changes of the brain perfusion in different diseases



Pick disease: atrophy in both frontal lobes



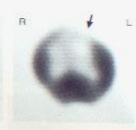
R R







Figure 2a



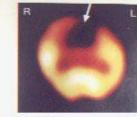










Figure 1a

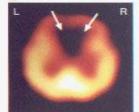


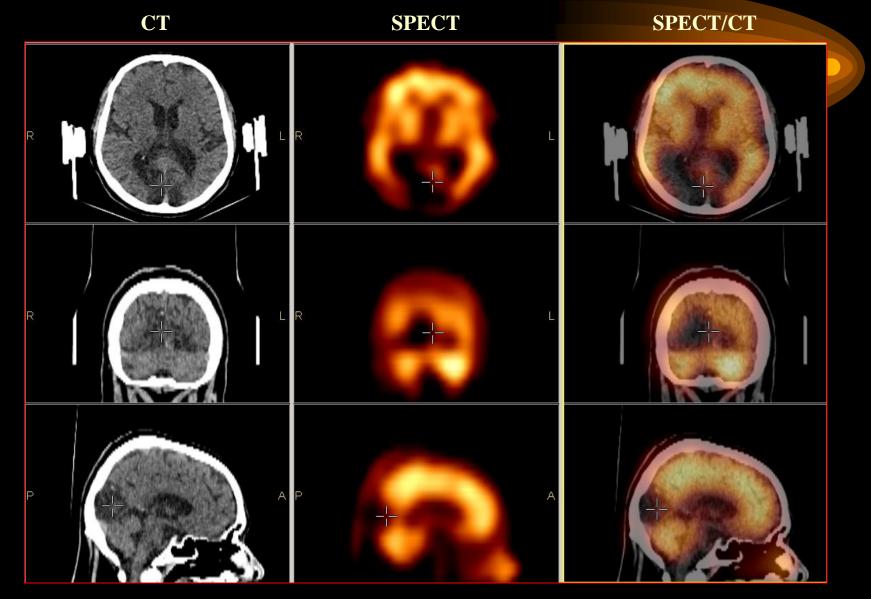
Figure 1b





R

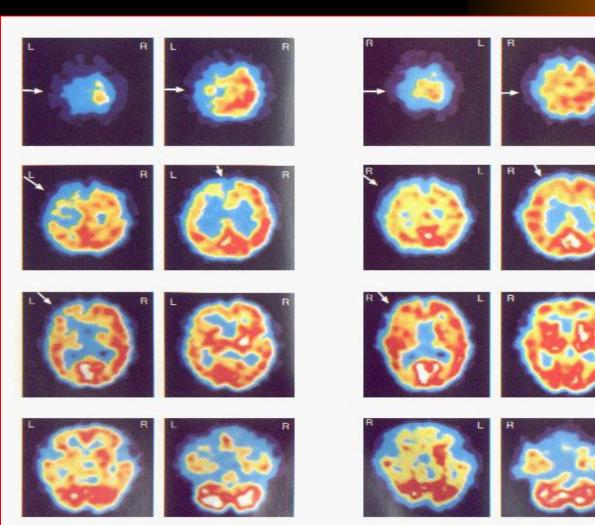
Perfusion defect (stroke) in the right occipital region (produced blindness)



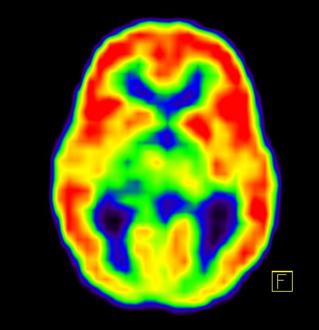
Occlusion of left internal carotis artery

Before operation

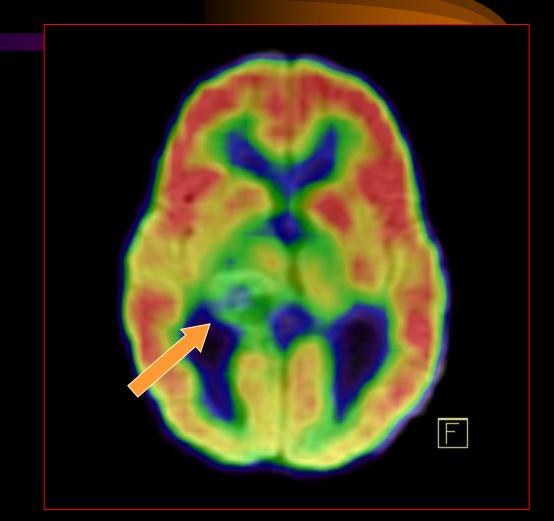
After operation



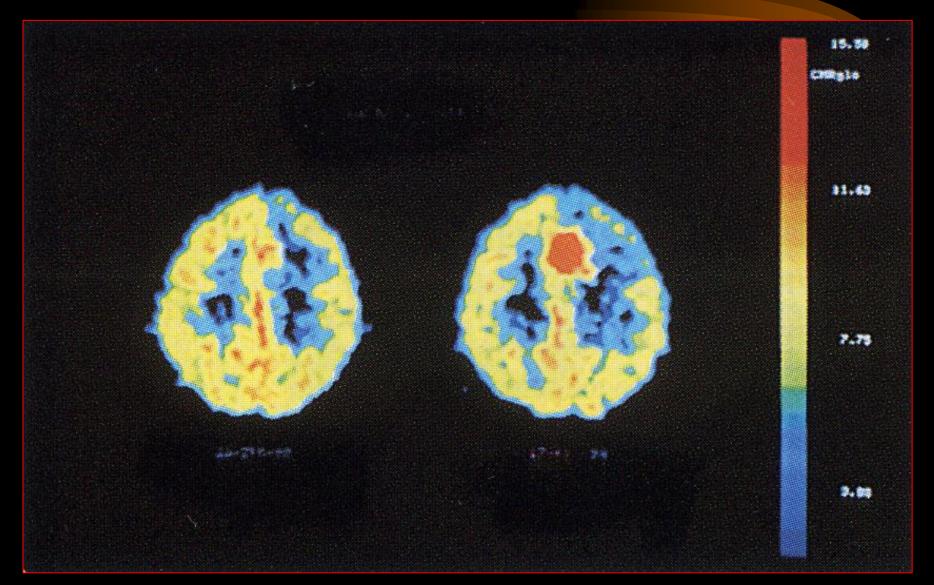




Cerebral glioma by 18-FDG-PET



Recidiv parasagittal meningeoma after operation (18-FDG)



Thank you for your attention!