

TRAITEMENT D'IMAGES MEDICALES

Illustration du rôle d'un ingénieur
au sein d'une structure hospitalière

Module « Bioingénierie, 8^o semestre, 2011
Ecole Centrale de Marseille.
Denis Mariano-Goulart
Faculté de médecine et CHRU de Montpellier

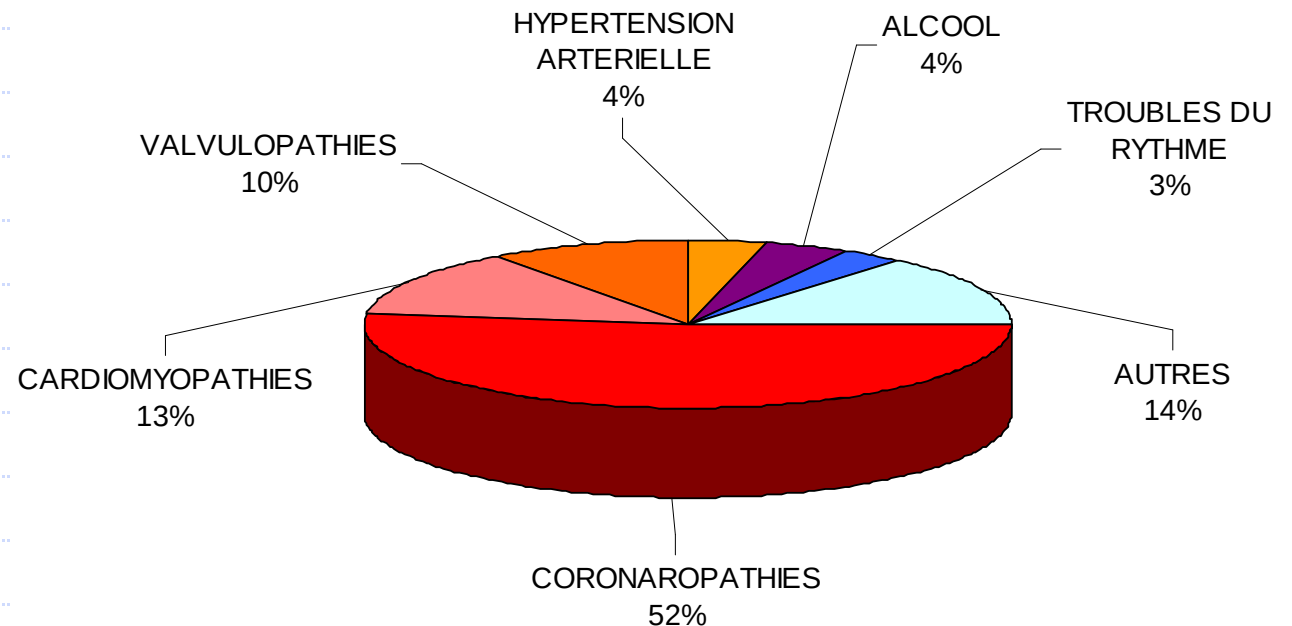
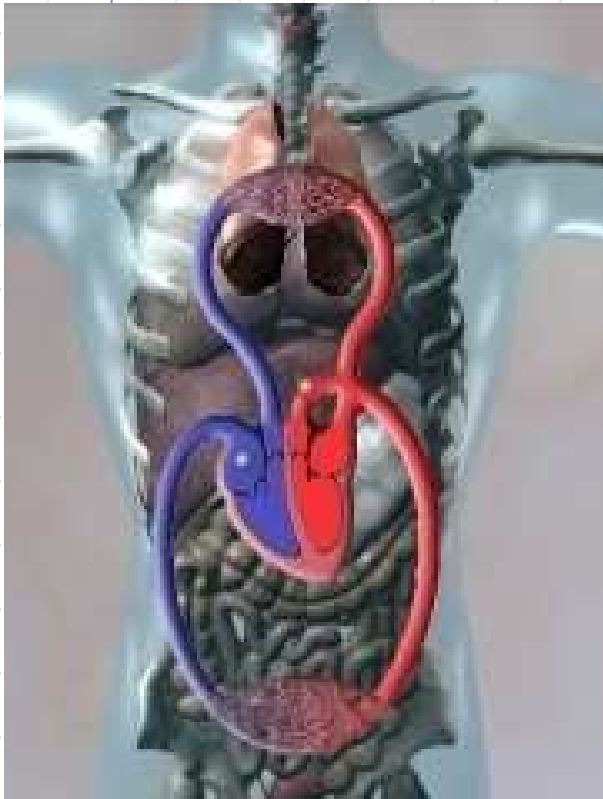
<http://scinti.etud.univ-montp1.fr>

PLAN

1. Problème médical à résoudre
2. Choix d'une modalité d'imagerie et d'un traceur
3. Acquisition des images
4. Reconstruction tomographique
5. Segmentation, analyse et validation (myocarde)
6. Segmentation, analyse et validation (ventriculo)

INSUFFISANCE CARDIAQUE

Insuffisance cardiaque : altération de la fonction pompe du coeur

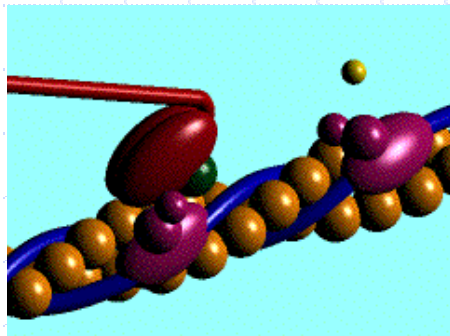
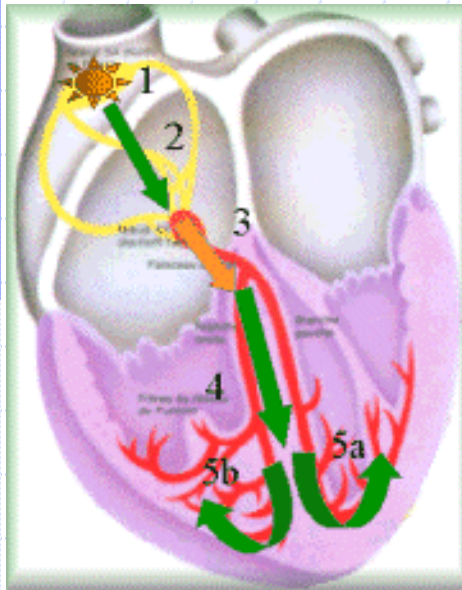


Prévalence = 1-3 % (1 million en France)

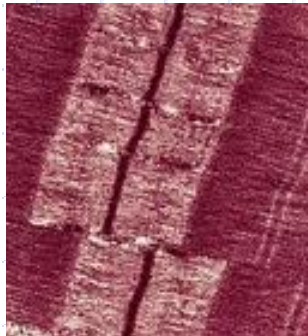
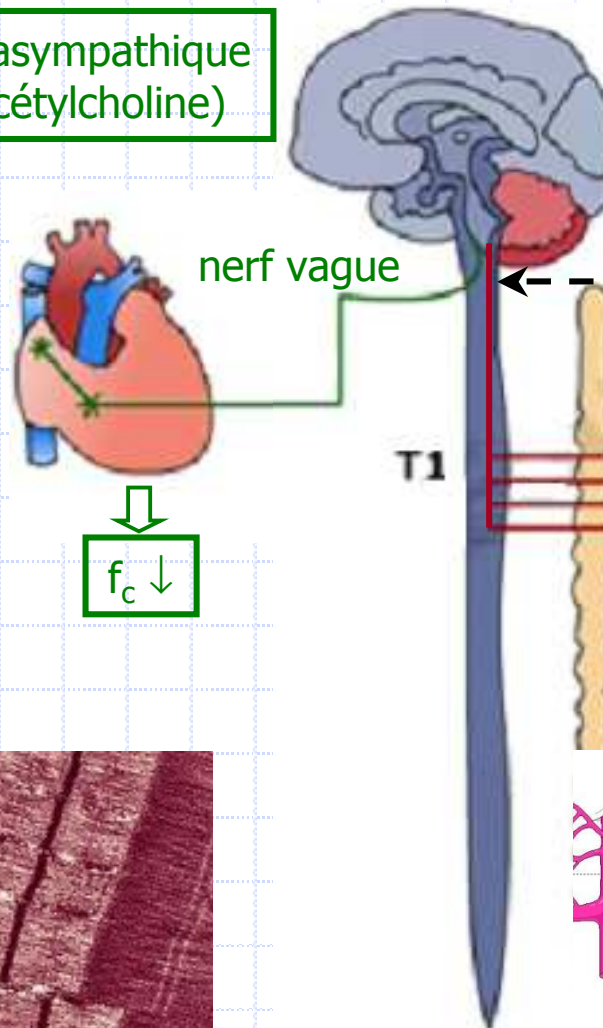
A 5 ans : 25 (H) à 40 %(F) de décès
Environ 1/2 morts subites

PHYSIOLOGIE CARDIAQUE

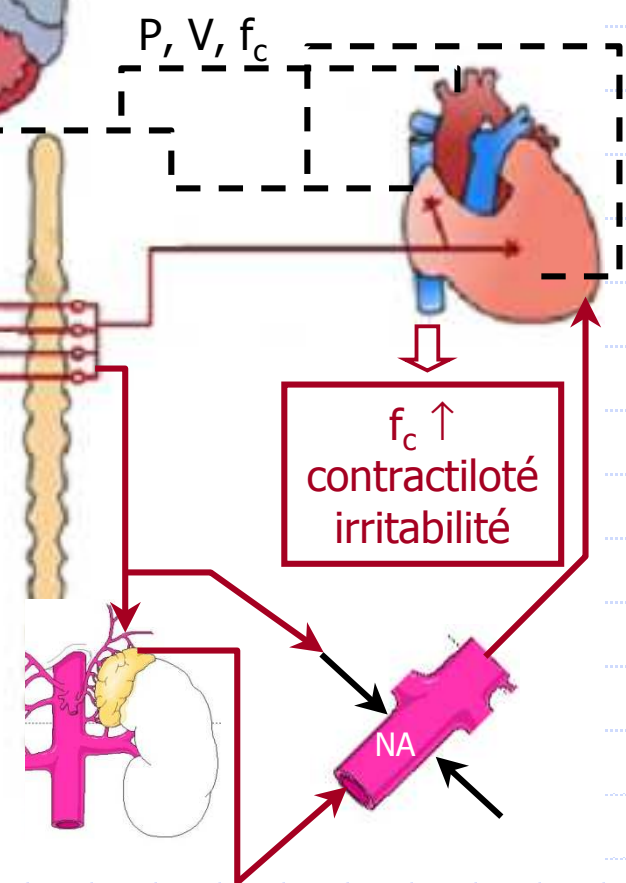
Automatisme
cardiaque



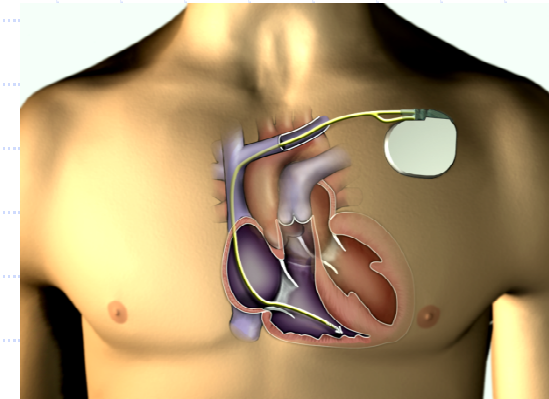
Parasympathique
(acétylcholine)



Sympathique
(noradrénaline)



PROBLEME CLINIQUE



- ◆ Mesures hygièno-diététiques

- ◆ Médicaments (Diu., IEC, β B)

- ◆ Défibrillateur implantable si

$$FE = \frac{VTD - VTS}{VTD} < 35\%$$

- ◆ mais décès avec $FE > 40\%$

- ◆ 35 % des défibrillateurs implantés sont sollicités / 3 ans.

- ◆ Meilleure identification des patients à risque de mort subite par imagerie ?

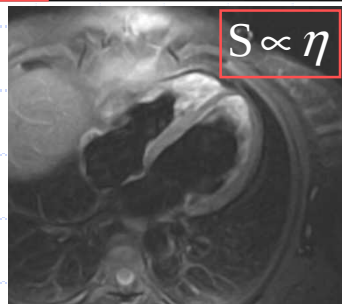
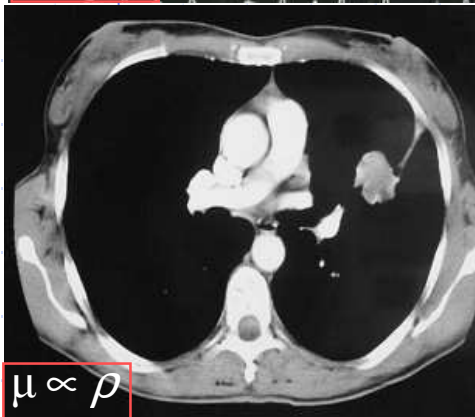
- ◆ Quelle fonction cibler (pompe, innervation, perfusion) ?

- ◆ Quelle modalité d'imagerie ?

- ◆ Quels traitements d'image ?

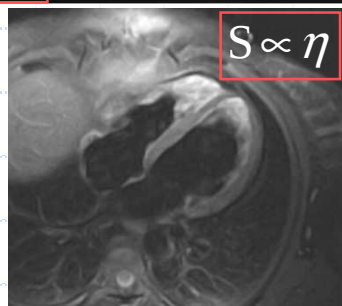
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CHOIX D'UNE MODALITE D'IMAGERIE



ANATOMIQUE

CHOIX D'UNE MODALITE D'IMAGERIE

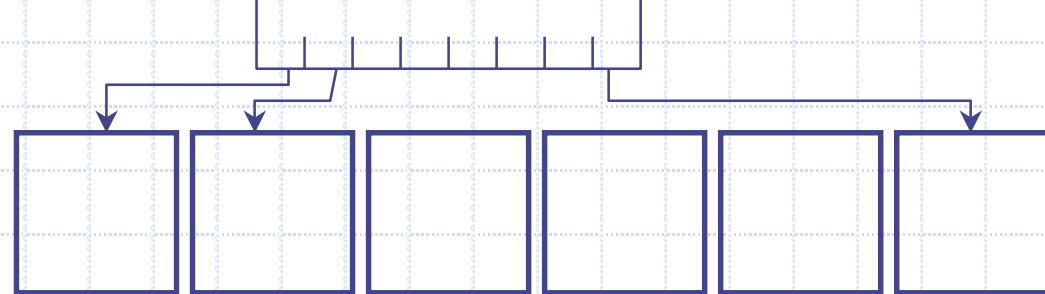
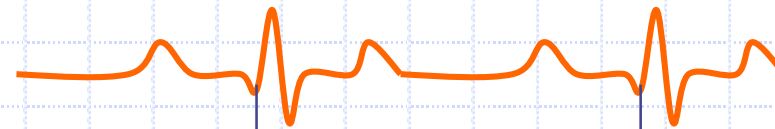


ANATOMIQUE

FONCTIONNELLE

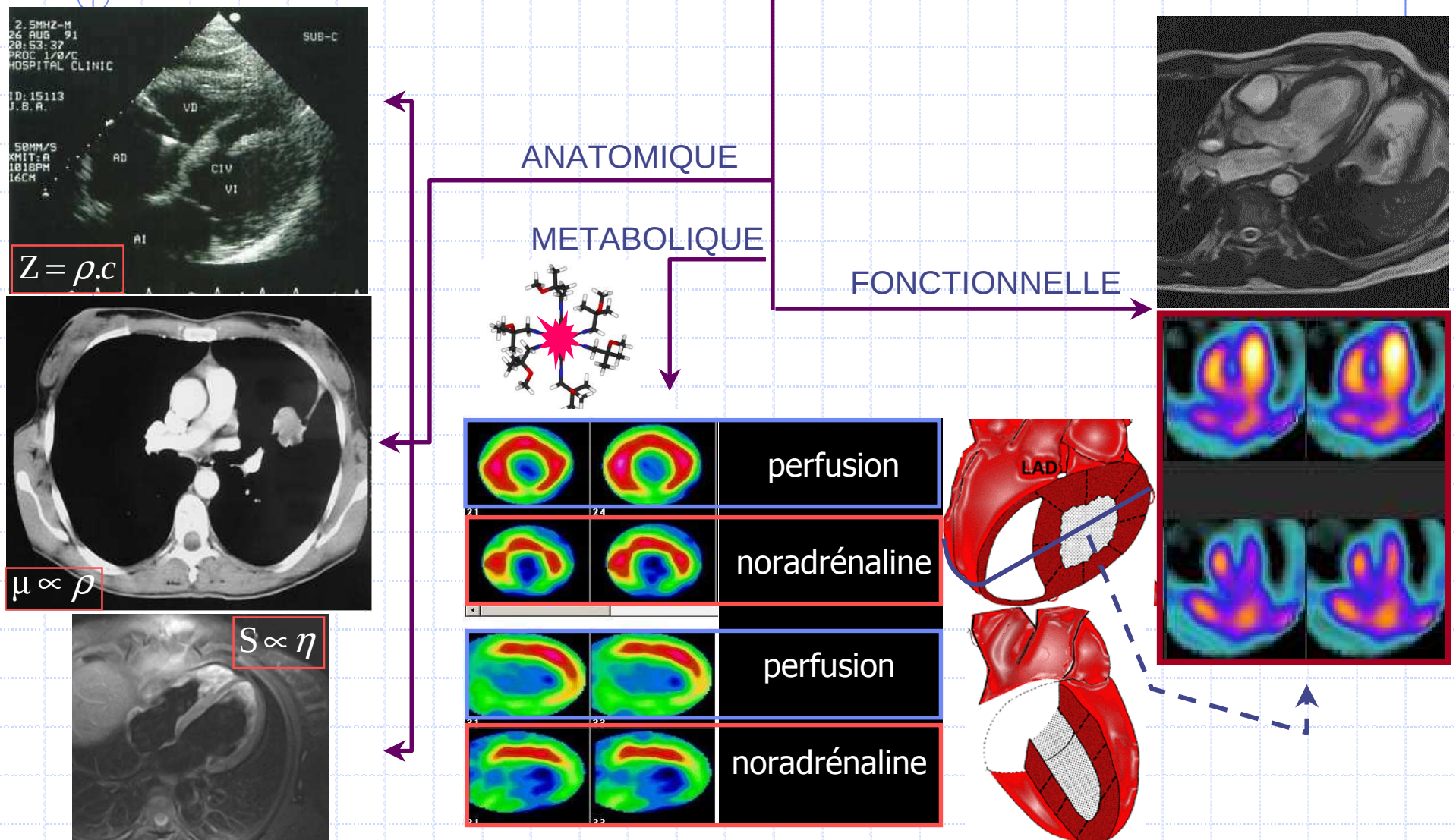


$$C = f(t)$$



Synchronisation à l'électrocardiogramme

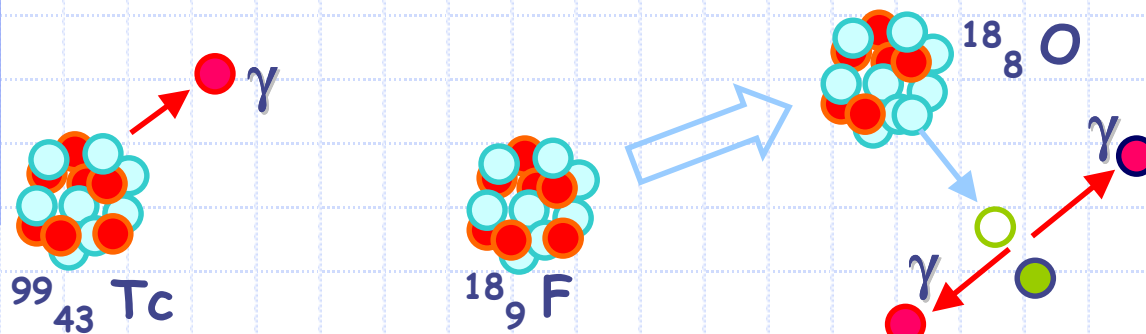
CHOIX D'UNE MODALITE D'IMAGERIE



CHOIX D'UN MARQUEUR RADIOACTIF

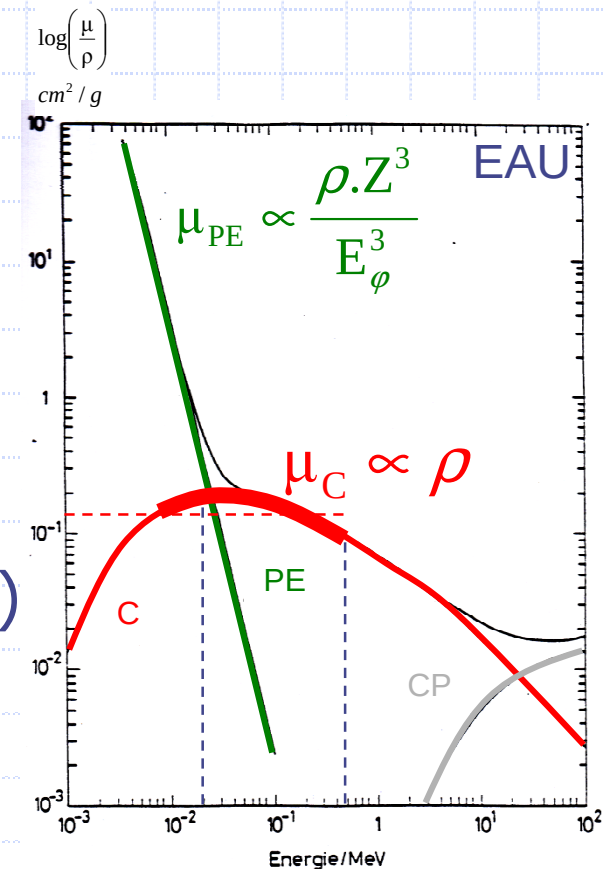
◆ Photons pénétrants et peu diffusés

- ◆ Émetteurs gamma ou bêta plus (ionisants)



◆ énergies détectables

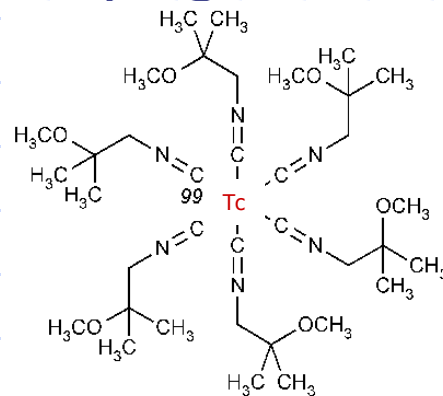
- ◆ γ (140 keV $^{99}_{43}\text{Tc}$ ou 159 keV $^{123}_{53}\text{I}$)
- ◆ $\beta^+ \rightarrow 2\gamma$ 511 keV ($^{18}_8\text{F}$)



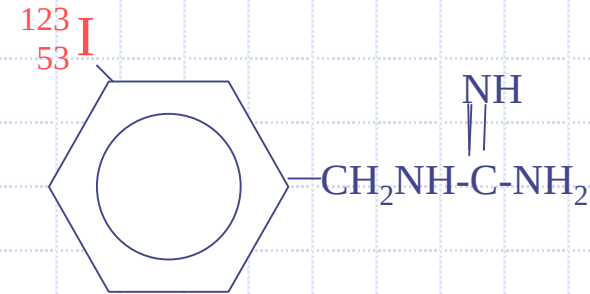
CHOIX D'UN RADIOTRACEUR

◆ Traçage possible d'un métabolisme

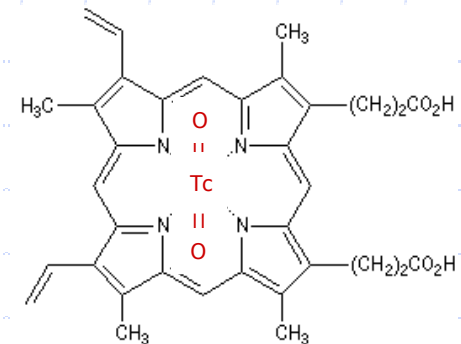
- ◆ Marquage de cations lipophiles (MIBI) au $^{99}_{43}\text{Tc}$



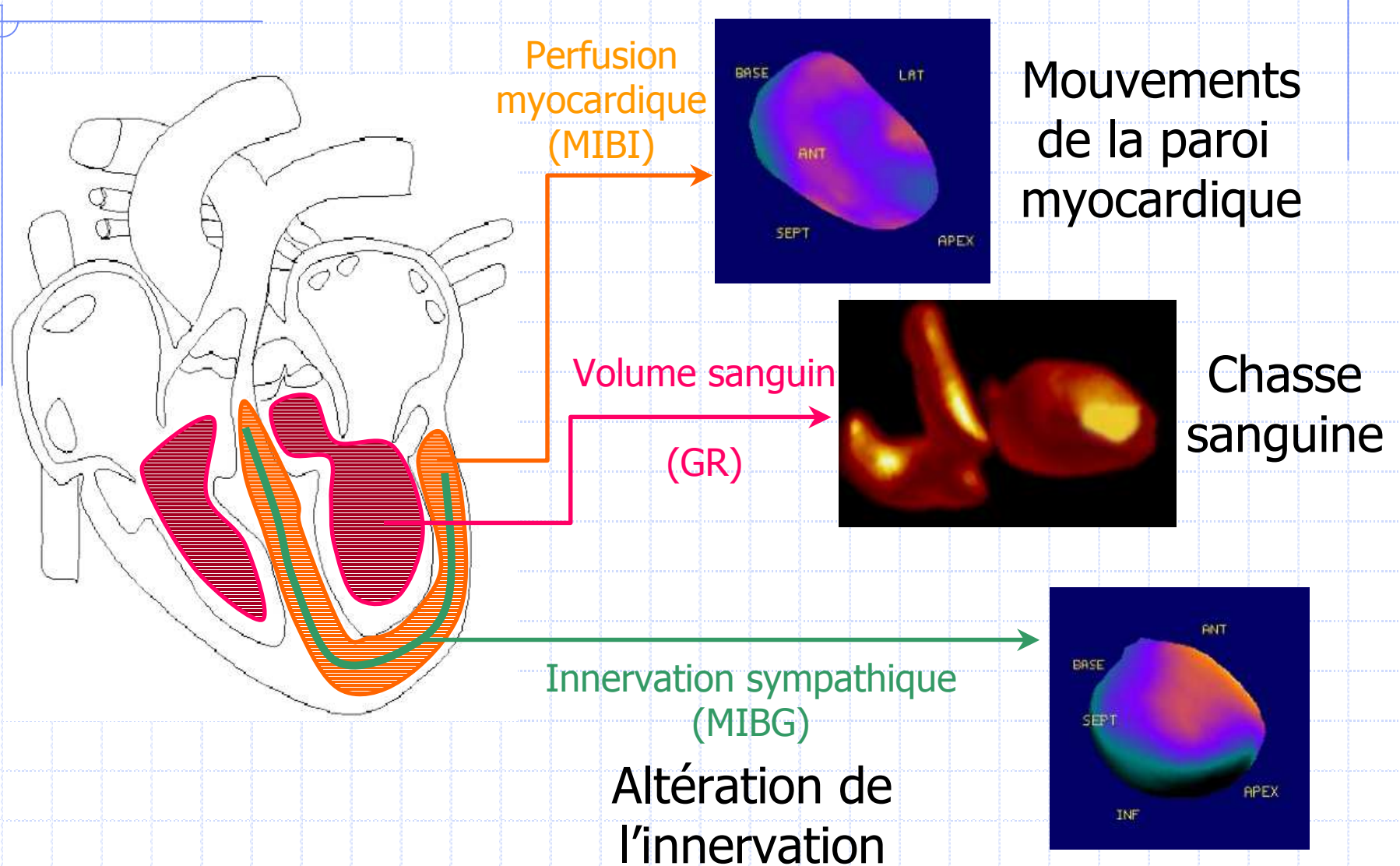
- ◆ Marquage de la MIBG à $^{123}_{53}\text{I}$



- ◆ Marquage des globules rouges au $^{99}_{43}\text{Tc}$
 - Chimie de complexation avec le noyau tétrapyrrolique de l'hème

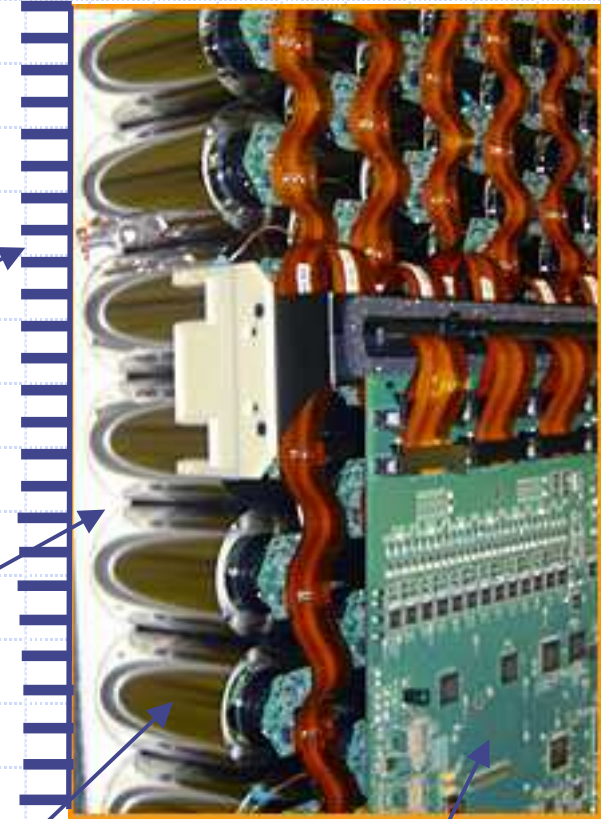
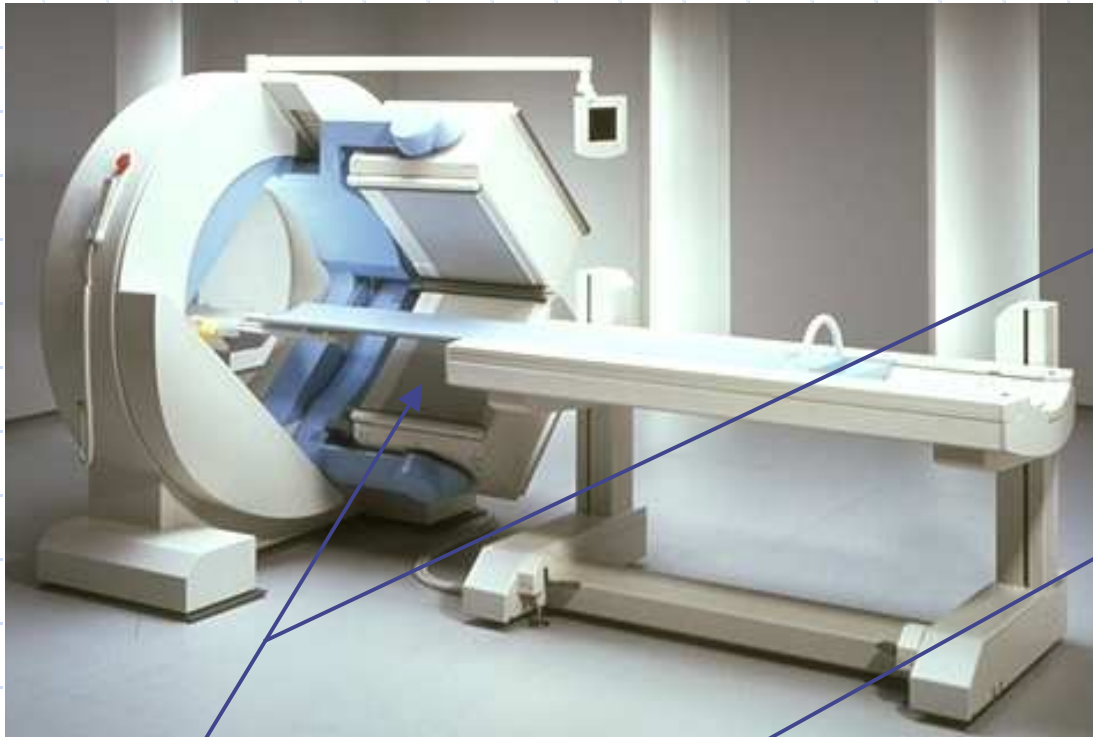


CHOIX D'UNE MODALITE D'IMAGERIE



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Single Photon Emission CT



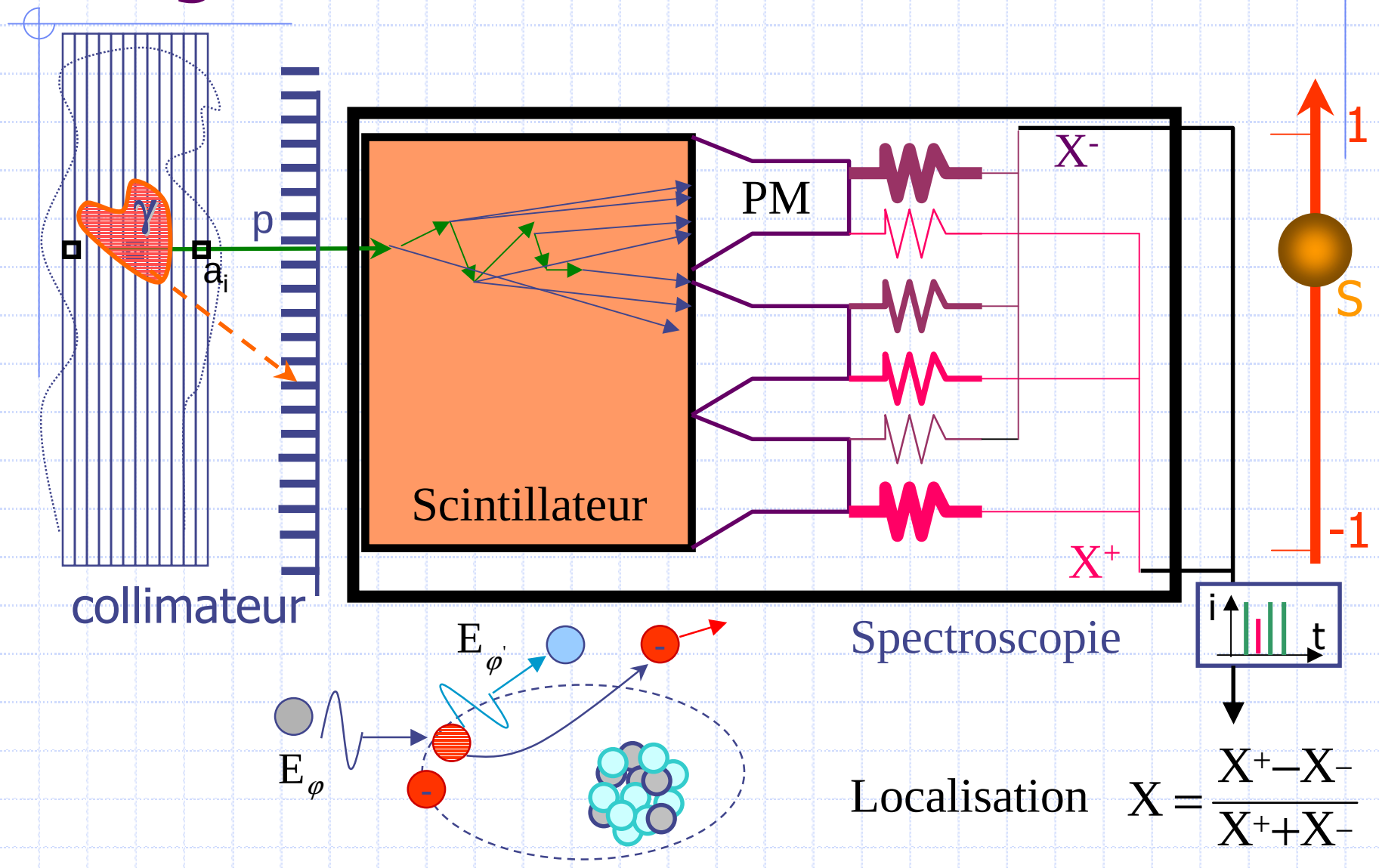
Collimateur

Scintillateur

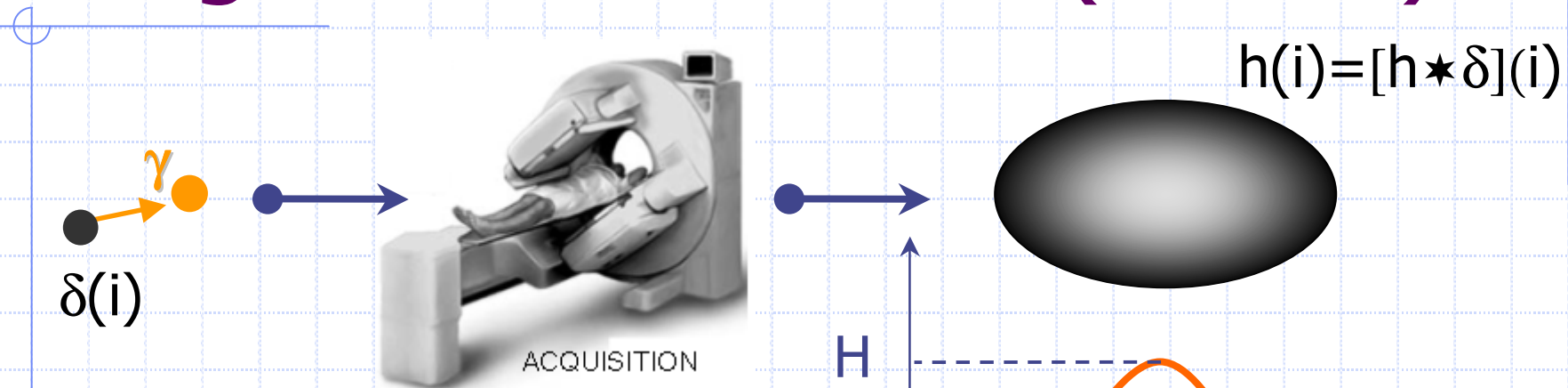
Photo-multiplicateur

Localisation

Single Photon Emission CT

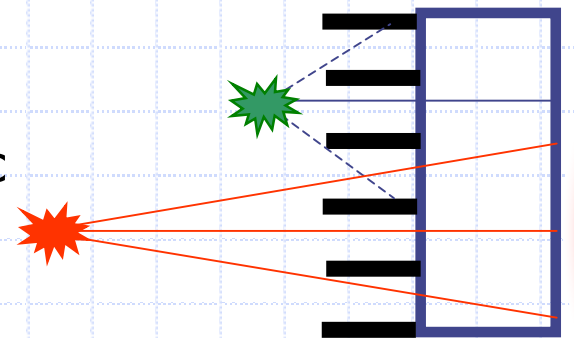


Largeur à mi-hauteur (FWHM)

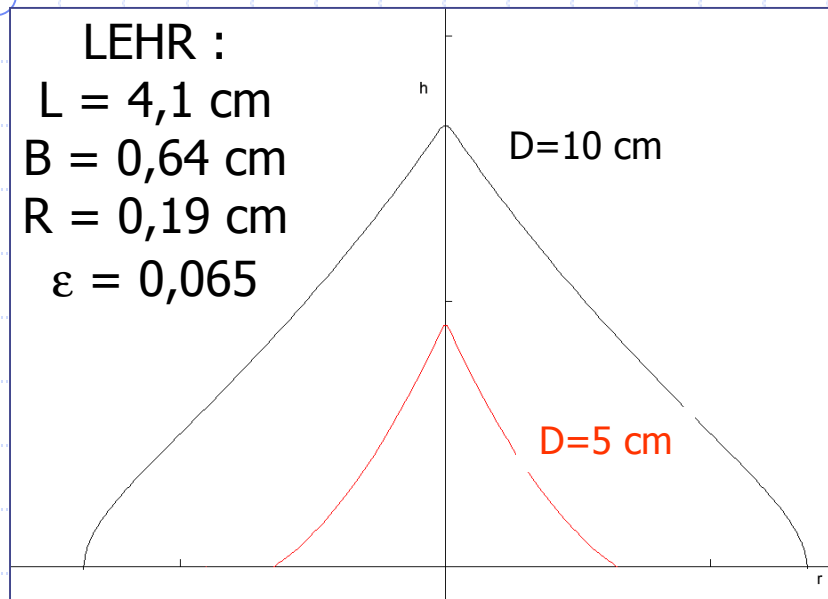


- Réponse intrinsèque h_i
 - ♦ $LMH \leq 4 \text{ mm}$, $\pm$ invariante

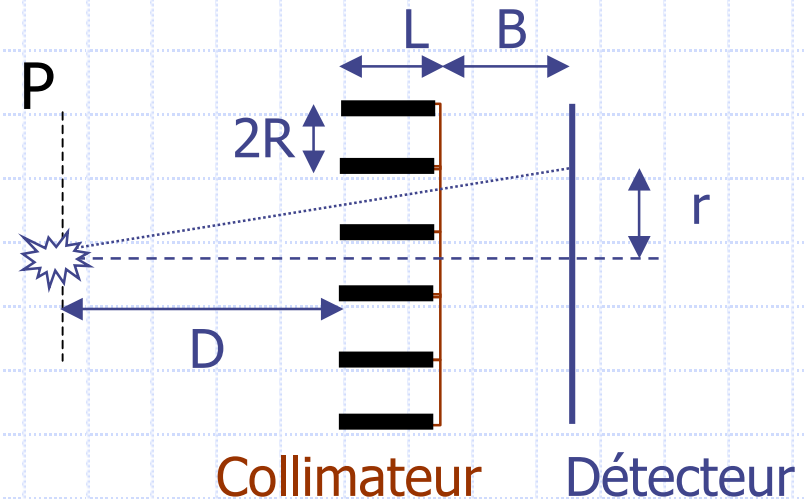
- Réponse du collimateur h_c
 - ♦ La LMH dépend de la distance source-détecteur



Réponse d'un collimateur



$$h(r) = \frac{\varepsilon}{\pi} \left[2 \cdot \arccos\left(\frac{\lambda \cdot r}{2}\right) - \lambda \cdot r \cdot \sqrt{1 - \left(\frac{\lambda \cdot r}{2}\right)^2} \right]$$

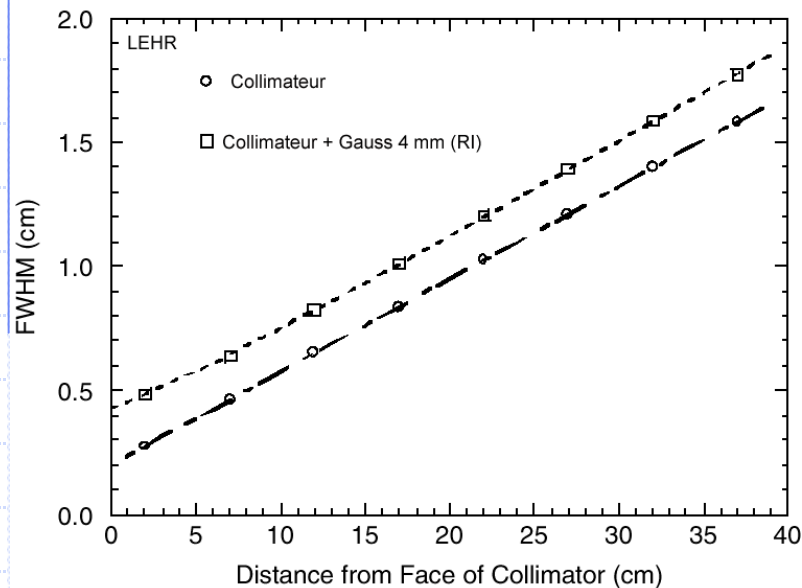


$$\lambda = \frac{L}{R \cdot (L + D + B)}$$

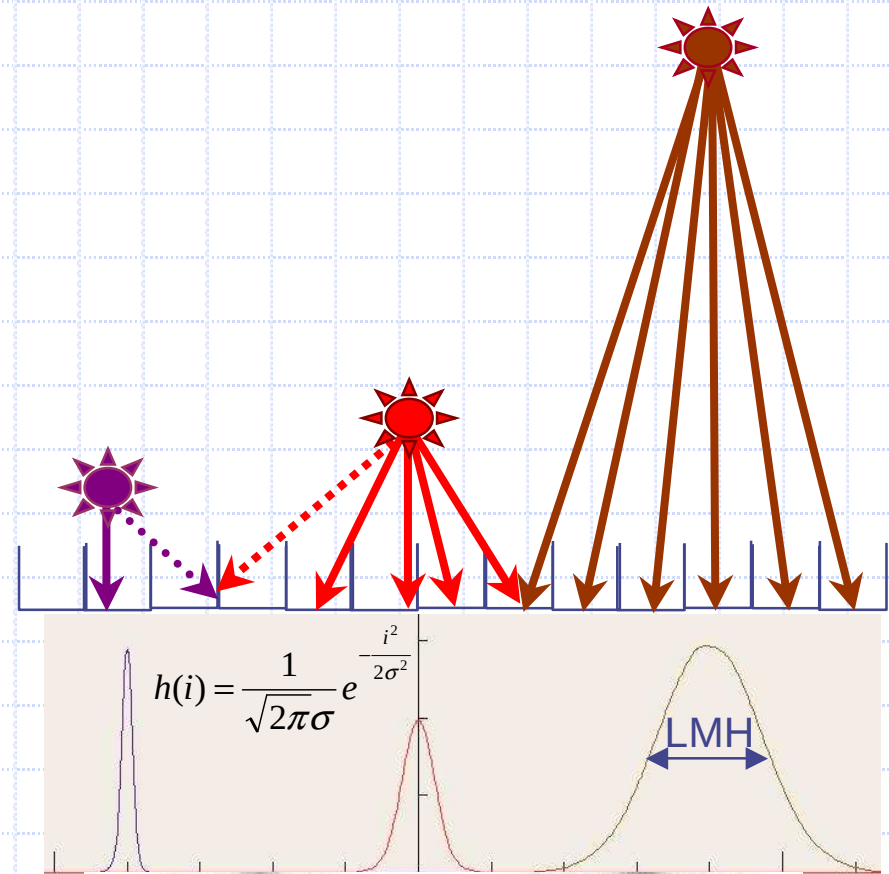
$$h_i \star h_c \approx \text{Gaussienne}$$

Linéarité entre LMH et D

$$h_i \star h_c \approx \text{Gaussienne}$$

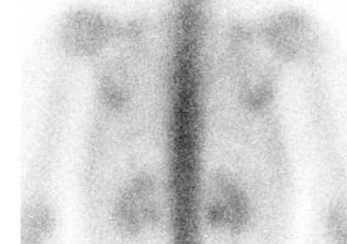


$$LMH = 2.\sqrt{2.\ln 2}\sigma = kD + k'$$



Utilité d'une déconvolution
(non stationnaire)

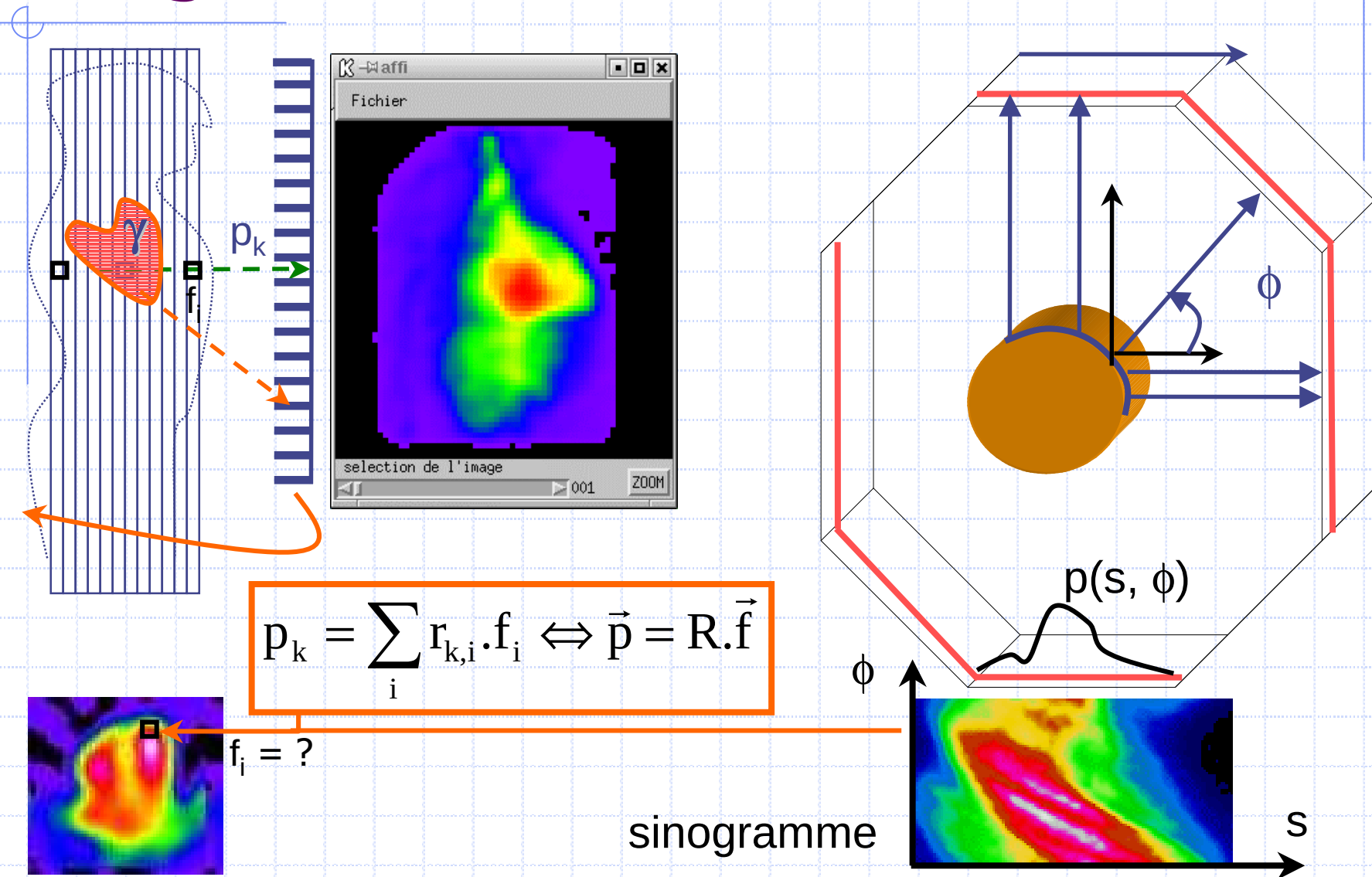
contact



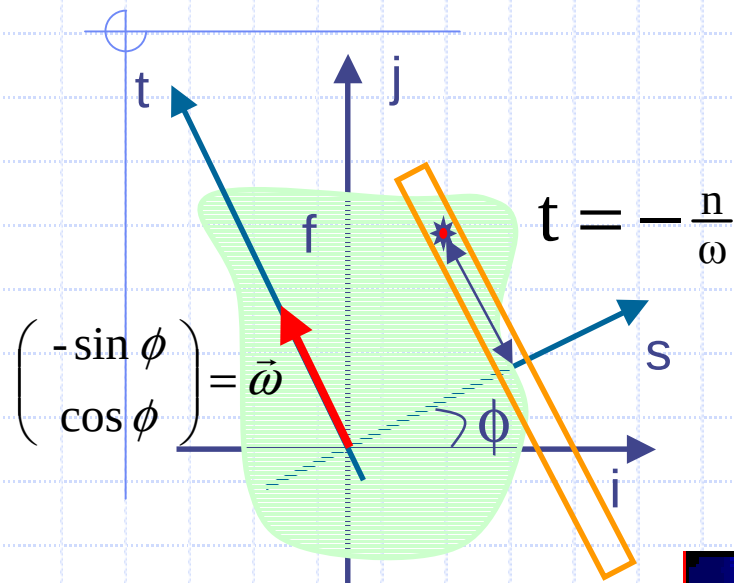
50 cm

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Single Photon Emission CT



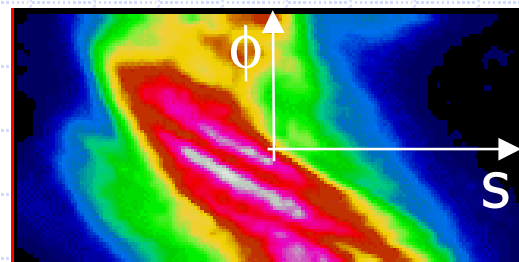
Déconvolution (hors synchro ECG)



sources à t mm du
centre de rotation

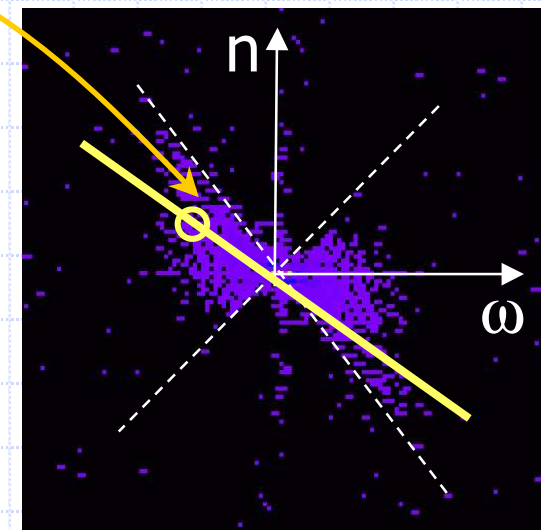
↓

signal $\approx$ sur la droite
 $n = -\omega \cdot t$ de la TF2
du sinogramme



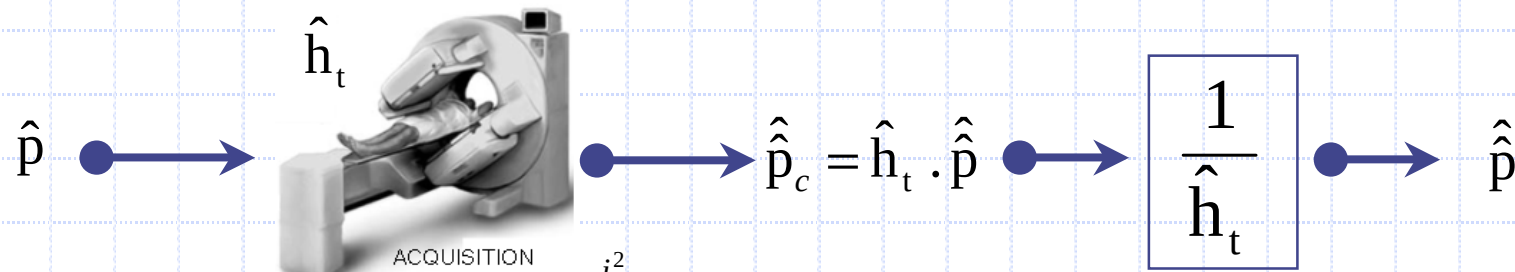
$$p_c(s, \phi) = \int_t f(s \vec{\omega}^\perp + t \vec{\omega}) dt$$

TF₂



$$\hat{p}_c(\sigma, n)$$

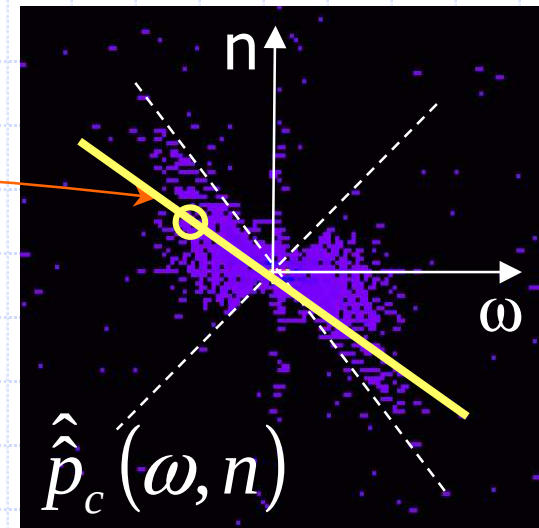
Principe fréquence-distance



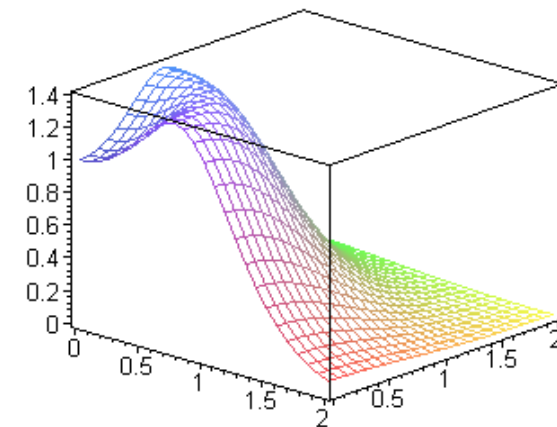
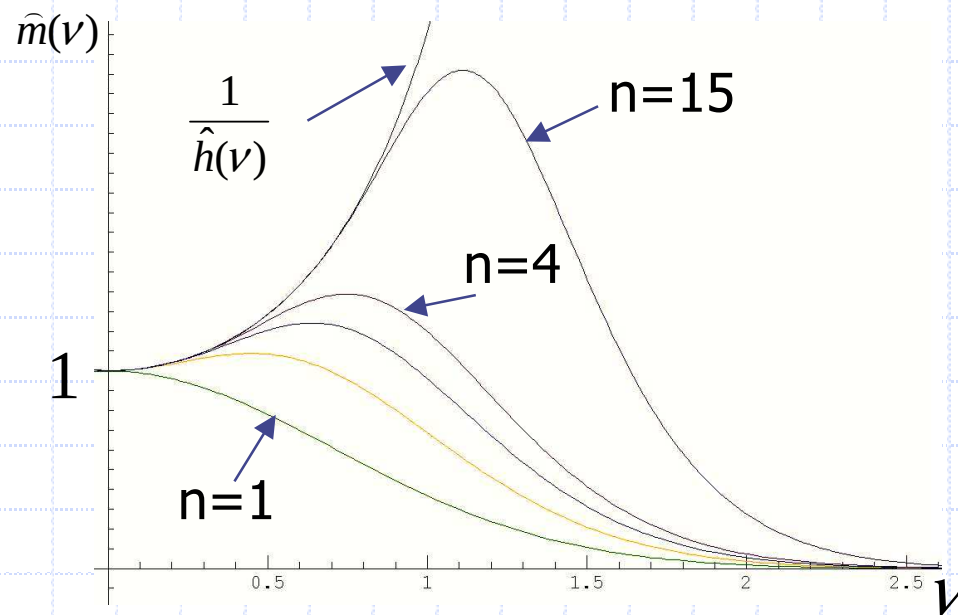
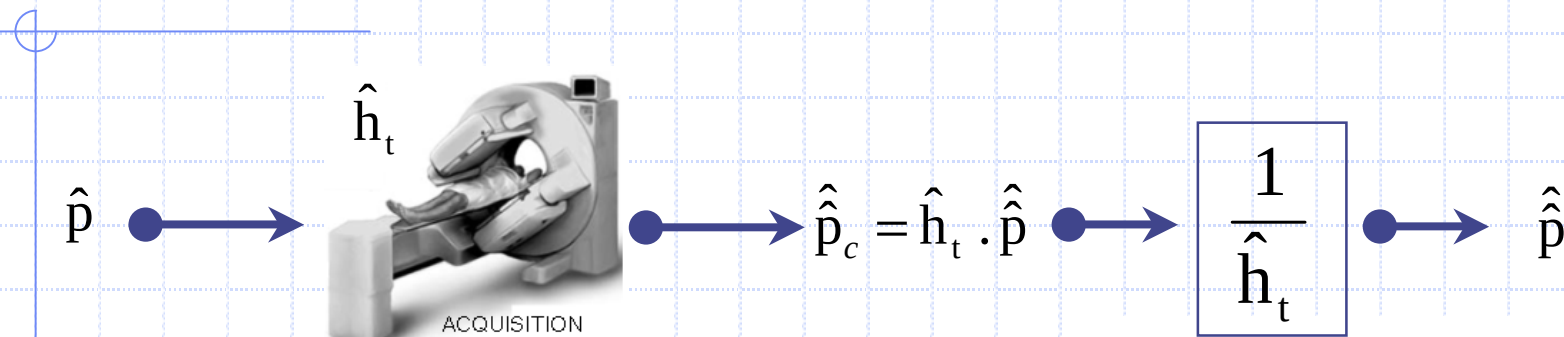
$$h_t(i) = \frac{1}{\sqrt{2\pi}\sigma_t} e^{-\frac{i^2}{2\sigma_t^2}}$$

$$\hat{p}_c(\omega, n) = \hat{h}_{\frac{n}{\omega}}(\omega) \cdot \hat{p}(\omega, n)$$

$$\hat{p}(\omega, n) = \frac{1}{\hat{h}_{\frac{n}{\omega}}(\omega)} \cdot \hat{p}_c(\omega, n)$$



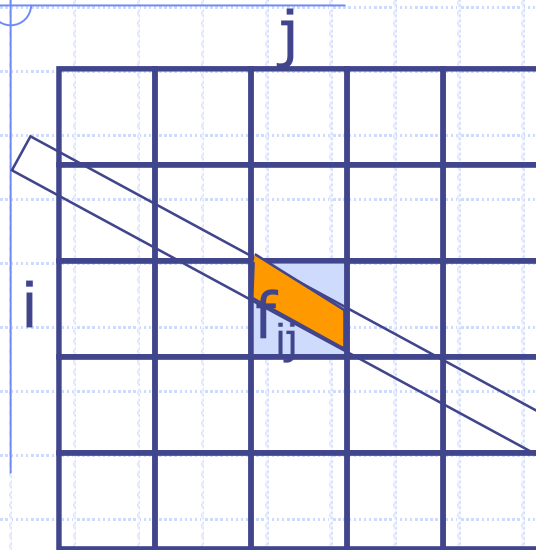
Filtre de déconvolution de Metz



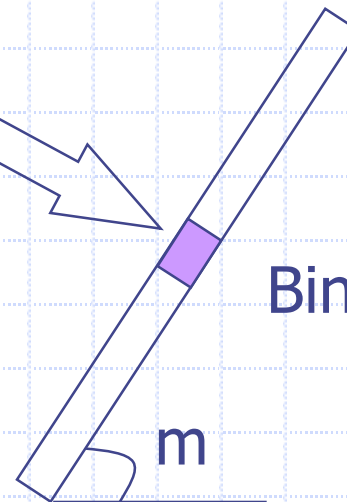
$$\hat{m}(v, v') = \frac{1 - \left[1 - \hat{h}(v, v')^2 \right]^n}{\hat{h}(v, v')}$$

$$n = 0,834 \cdot \ln(C) - 7,774$$

Alternative pour déconvoluer

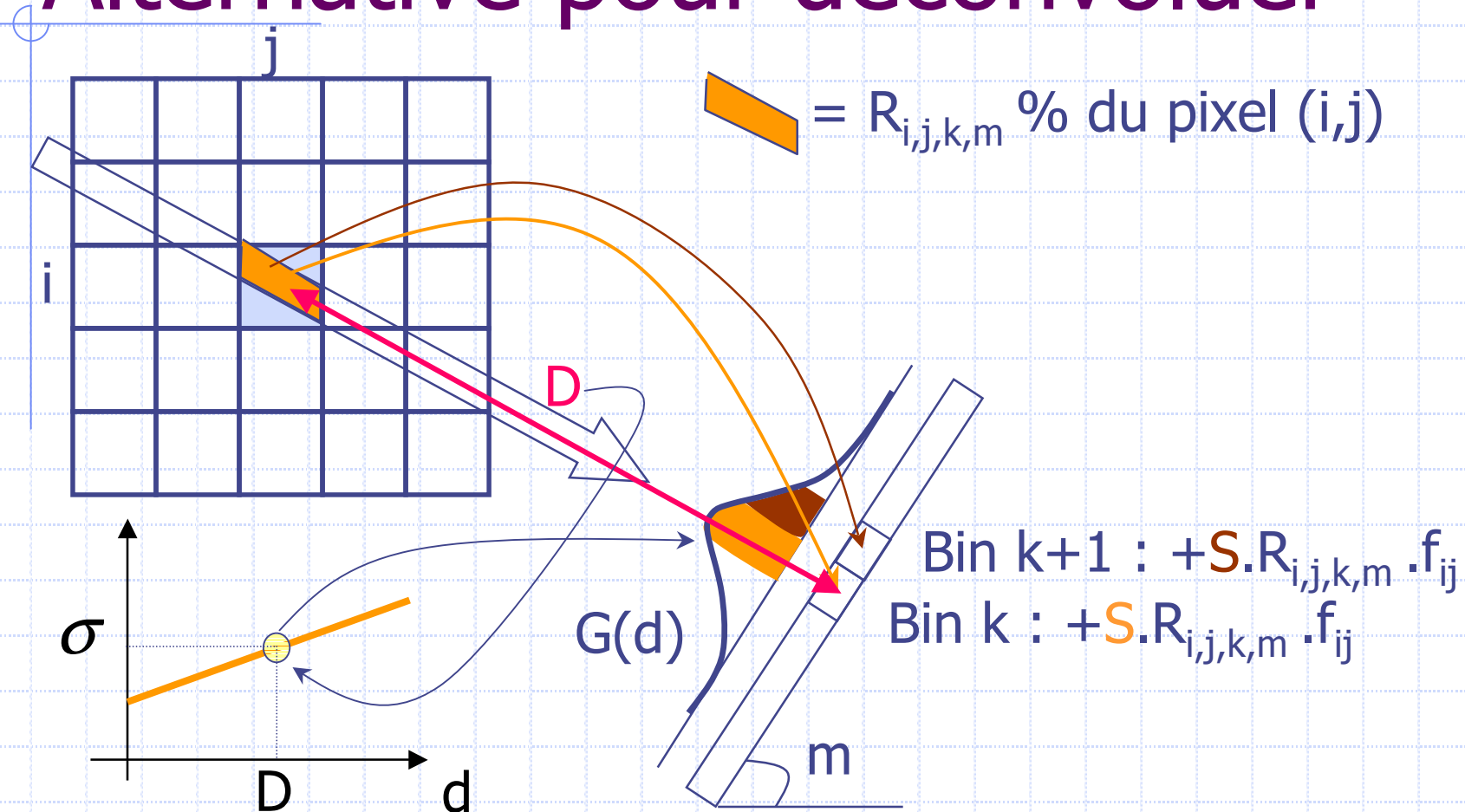


 = $R_{i,j,k,m}$ % du pixel (i,j)



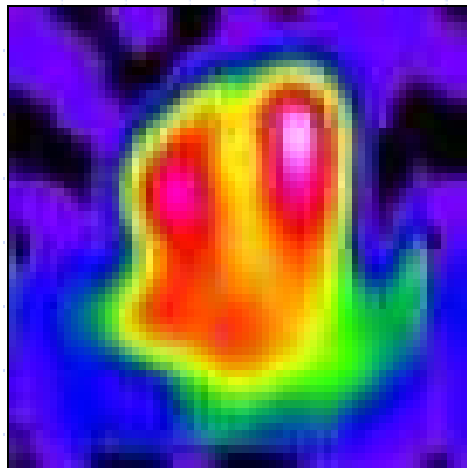
Bin k : $+ R_{i,j,k,m} \cdot f_{ij}$

Alternative pour déconvoluer

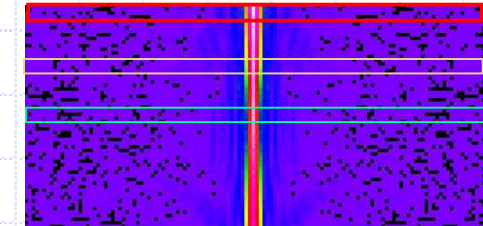
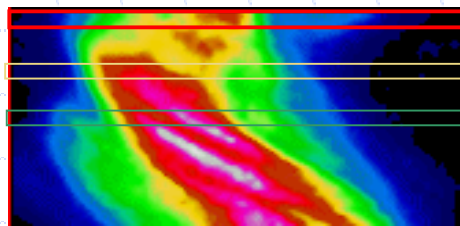
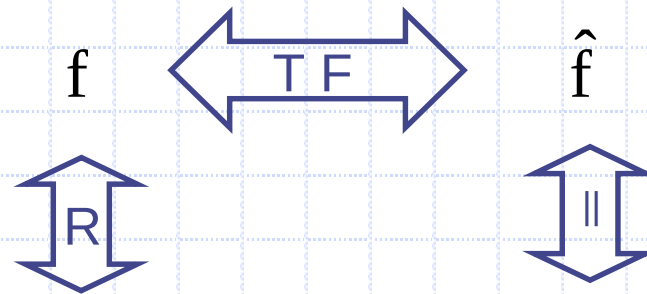
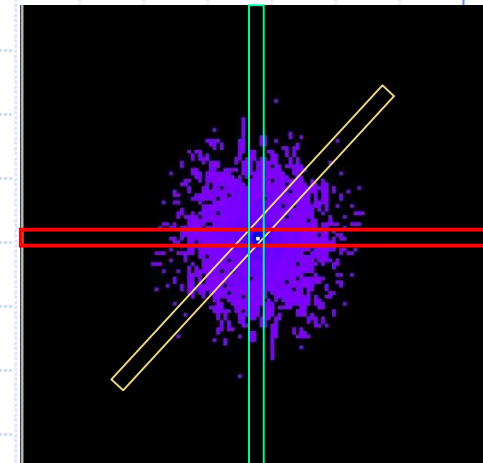


Solvabilité du problème tomographique

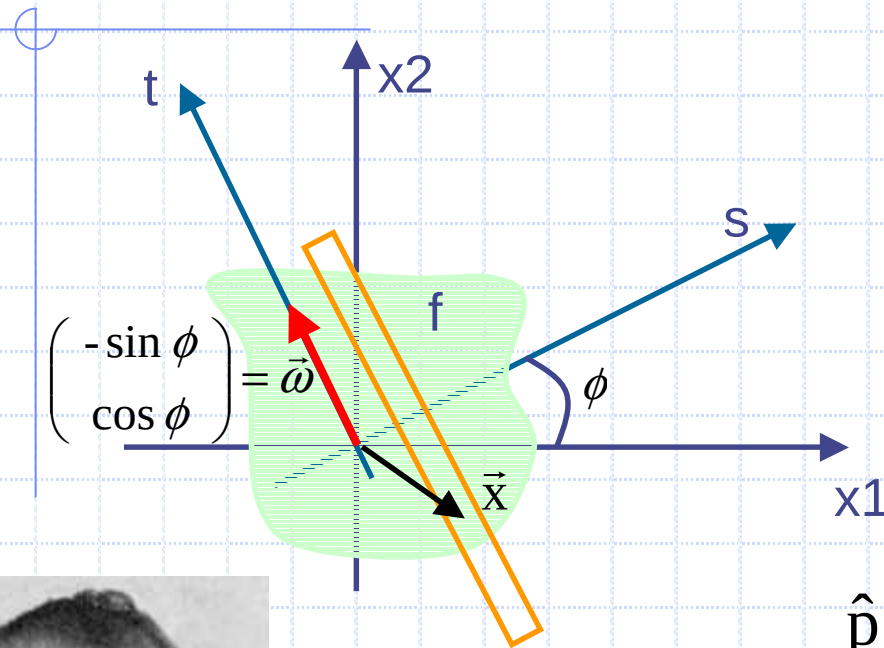
Aspect analytique: R bijectif d'inverse continu



$$\hat{p}_{\vec{\omega}^\perp}(\sigma) = \hat{f}(\sigma \cdot \vec{\omega}^\perp)$$



Démonstration



$$\begin{cases} p_{\vec{\omega}}(s) = \int_t f(s\vec{\omega}^\perp + t\vec{\omega}) dt \\ \hat{p}_{\vec{\omega}}(\sigma) = \int_s p_{\vec{\omega}}(s) \cdot e^{-i.s.\sigma} ds \end{cases}$$

$$\hat{p}_{\vec{\omega}}(\sigma) = \int_s \int_t f(s\vec{\omega}^\perp + t\vec{\omega}) e^{-i.s.\sigma} dt ds$$

$$\hat{p}_{\vec{\omega}}(\sigma) = \iint f(\vec{x}) e^{-i.\sigma \vec{x}.\vec{\omega}^\perp} d\vec{x}$$

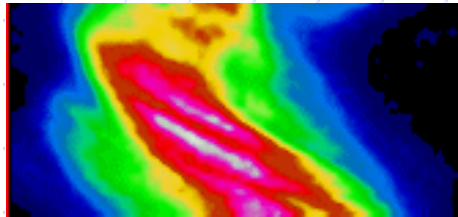
$$\hat{p}_{\vec{\omega}}(\sigma) = \hat{f}(\sigma.\cos \phi, \sigma.\sin \phi) = \hat{f}(\sigma.\vec{\omega}^\perp)$$



J. Radon

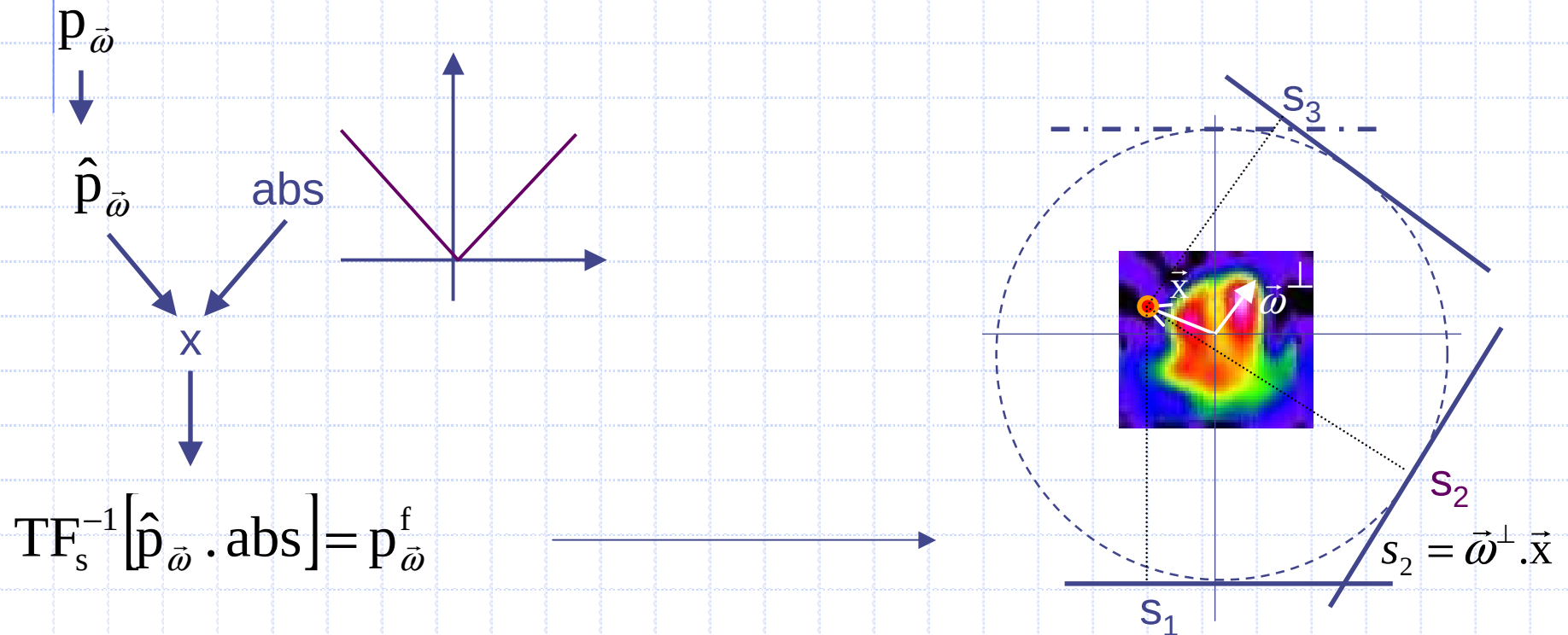
1887-1956

Rétroprojection filtrée



$$f(\vec{x}) = (R^* p^f)(\vec{x})$$

Projections sur 180°



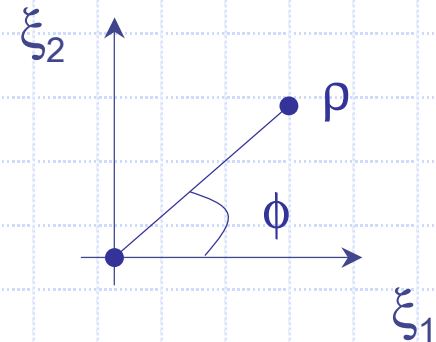
Démonstration

$$f(\vec{x}) = \iint \hat{f}(\vec{\xi}) e^{i\vec{x} \cdot \vec{\xi}} d\vec{\xi}$$

$$f(\vec{x}) = \int_{\phi=0}^{\pi} \int_{\sigma=-\infty}^{+\infty} \hat{f}(\sigma \vec{\omega}^{\perp}) e^{i\sigma \vec{\omega}^{\perp} \cdot \vec{x}} |\sigma| d\sigma d\phi$$

$$f(\vec{x}) = \int_{\phi=0}^{\pi} \int_{\sigma=-\infty}^{+\infty} \hat{p}_{\vec{\omega}^{\perp}}(\sigma) |\sigma| e^{i\sigma \vec{\omega}^{\perp} \cdot \vec{x}} d\sigma d\phi$$

$$\underbrace{\text{TF}_s^{-1}[\hat{p}_{\vec{\omega}^{\perp}} \cdot \text{abs}]}_{p_{\vec{\omega}}^f}(\vec{\omega}^{\perp} \cdot \vec{x})$$



J. Radon
1887-1956

$$f(\vec{x}) = (R^* p^f)(\vec{x})$$

Solvabilité du problème tomographique

Aspect discret : le problème est moins simple !

✓ R surjectif ? :

$${}^tR.R\vec{f} = {}^tR.\vec{p} \Leftarrow \bar{f} = \arg \min_f \|\vec{p} - R\vec{f}\|^2$$

✓ R injectif ? :

✓ Norme de R^{-1} ?

R^{-1} continue mais $\|R^{-1}\|$ grande

← choix parmi les solutions possibles

$$\frac{\|\delta\vec{f}\|}{\|\vec{f}\|} \leq \kappa(R) \frac{\|\delta\vec{p}\|}{\|\vec{p}\|} \quad \text{où} \quad \kappa(R) = \|R\| \|R^{-1}\| = \frac{\lambda_{\max}}{\lambda_{\min}} \gg 1$$

Gradient Conjugué

$$\bar{f} = \arg \min_{f \in C} \| R\vec{f} - \vec{p} \|^2$$

Initialisation :

$$\vec{d}^0 = \vec{r}^0 = R^* \cdot \vec{p} \longrightarrow$$

$$\omega^j = \frac{\|\vec{r}^j\|^2}{\langle \vec{d}^j | R^* \cdot R \cdot \vec{d}^j \rangle}$$

$$\vec{r}^{j+1} = \vec{r}^j - \omega^j \cdot R^* \cdot R \cdot \vec{d}^j$$

$$\vec{d}^{j+1} = \vec{r}^{j+1} + \frac{\|\vec{r}^{j+1}\|^2}{\|\vec{r}^j\|^2} \cdot \vec{d}^j$$

$$\vec{f}^{j+1} = \vec{f}^j + \omega^j \cdot \vec{d}^j$$

Estimation de $\kappa(R)$ par $Rf=p$

◆ Gradient conjugué

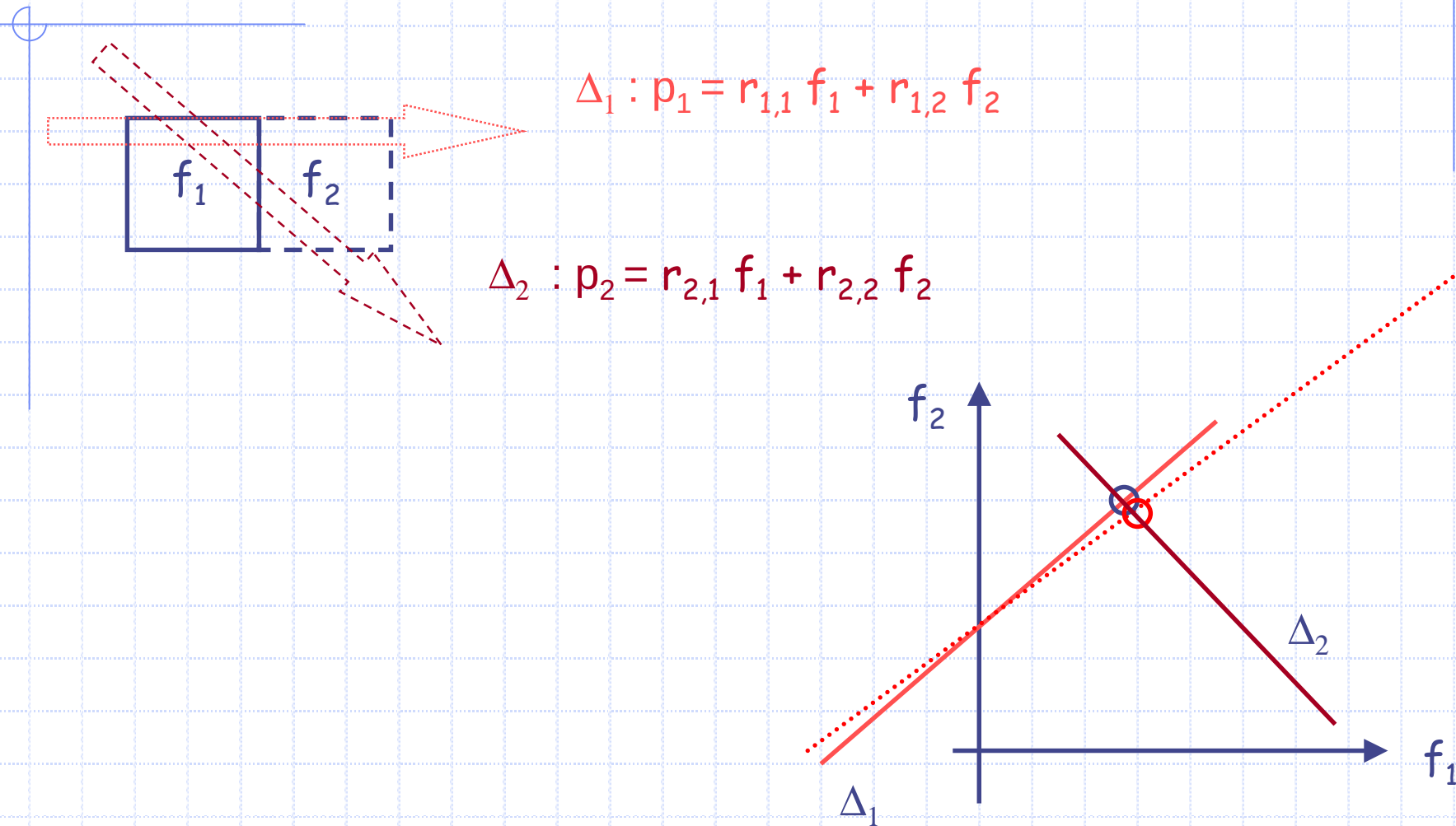
◆ Approximation de Galerkin

$$\omega^j = \frac{\|\vec{r}^j\|^2}{\langle \vec{d}^j | R^* . R . \vec{d}^j \rangle} , \quad \beta^j = \frac{\|\vec{r}^{j+1}\|^2}{\|\vec{r}^j\|^2}$$

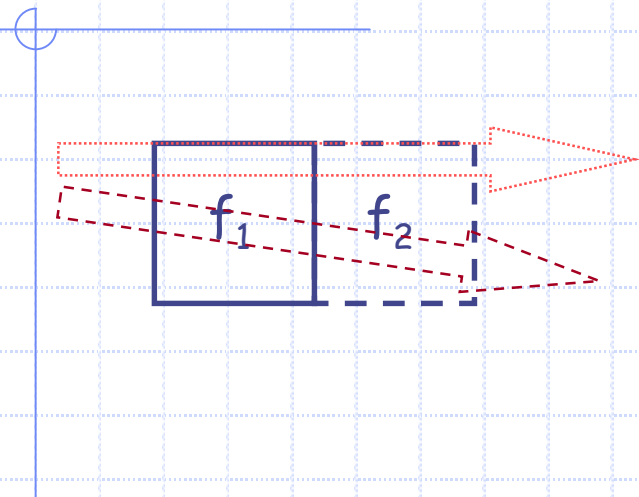
$$\begin{pmatrix} \frac{1}{\omega^0} & -\frac{\sqrt{\beta^0}}{\omega^0} & 0 & 0 \\ -\frac{\sqrt{\beta^0}}{\omega^0} & \frac{1}{\omega^1} + \frac{\beta^0}{\omega^0} & \ddots & 0 \\ 0 & \ddots & \ddots & -\frac{\sqrt{\beta^{j-1}}}{\omega^{j-1}} \\ 0 & 0 & -\frac{\sqrt{\beta^{j-1}}}{\omega^{j-1}} & \frac{1}{\omega^j} + \frac{\beta^{j-1}}{\omega^{j-1}} \end{pmatrix}$$

- ◆ dont le spectre CV vers celui de R : estimation de $\kappa(R)$
 - $\kappa(R)$ qui borne l'erreur sur la solution et le spectre

Approche intuitive (I)



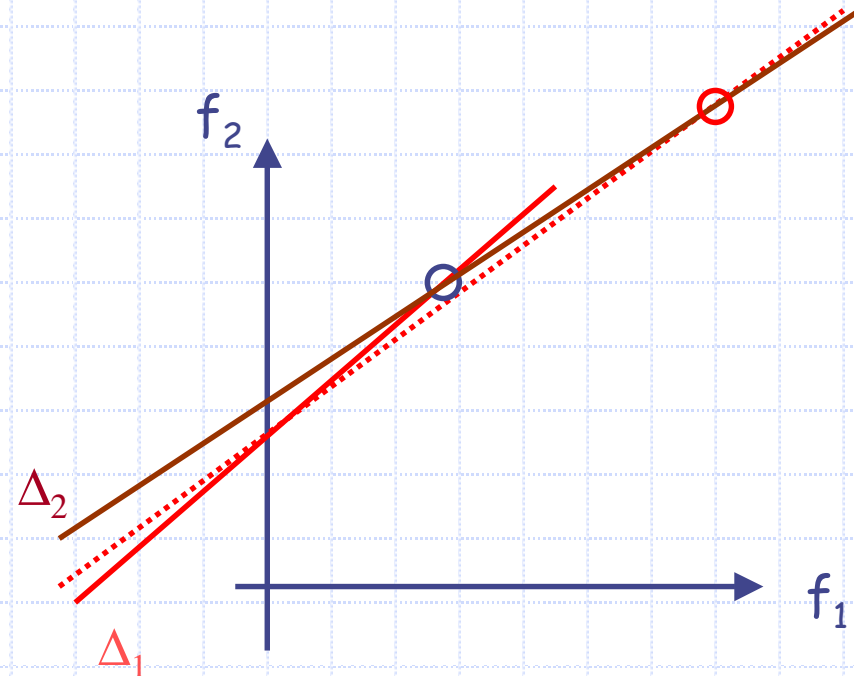
Approche intuitive (II)



$$\Delta_1 : p_1 = r_{1,1} f_1 + r_{1,2} f_2$$

$$\Delta_2 : p_2 = r_{2,1} f_1 + r_{2,2} f_2$$

$$\begin{aligned} 64^2 &= 4\,096 \\ 128^2 &= 16\,384 \\ 256^2 &= 65\,536 \\ 512^2 &= 262\,144 \end{aligned}$$



Exemple

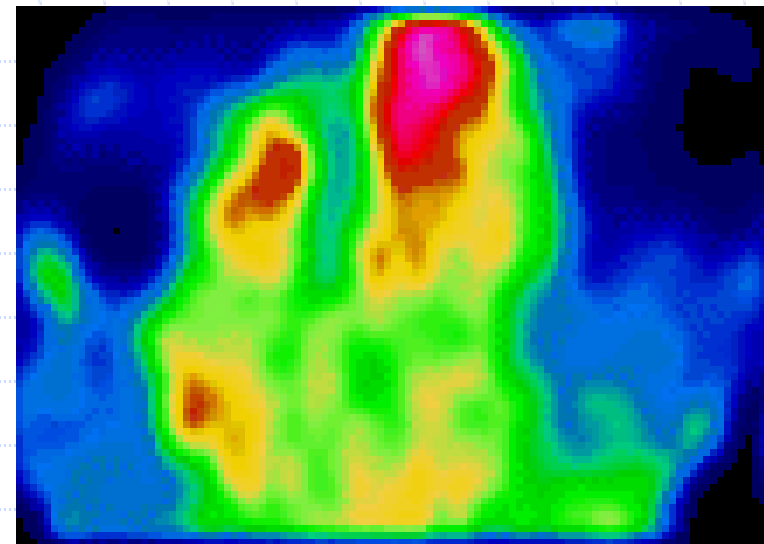
Matrice de Wilson

$$\begin{pmatrix} 10 & 7 & 8 & 7 \\ 7 & 5 & 6 & 5 \\ 8 & 6 & 10 & 9 \\ 7 & 5 & 9 & 10 \end{pmatrix} \begin{pmatrix} 1 \\ 1 \\ 1 \\ 1 \end{pmatrix} = \begin{pmatrix} 32 \\ 23 \\ 33 \\ 31 \end{pmatrix}$$

$$\text{Sp}(R) \approx \{0,010; 0,843; 3,858; 30,289\} \Rightarrow \kappa(R) = \frac{30,289}{0,010} = 2984$$

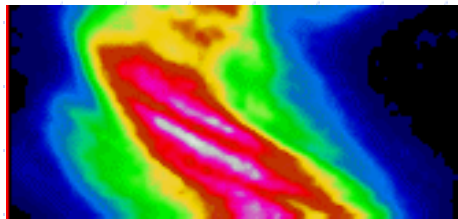
$$\begin{pmatrix} 10 & 7 & 8 & 7 \\ 7 & 5 & 6 & 5 \\ 8 & 6 & 10 & 9 \\ 7 & 5 & 9 & 10 \end{pmatrix} \begin{pmatrix} 9,2 \\ -12,6 \\ 4,5 \\ -1,1 \end{pmatrix} = \begin{pmatrix} 32,1 \\ 22,9 \\ 33,1 \\ 30,9 \end{pmatrix}$$

$$\begin{pmatrix} 10 & 7 & 8 & 7 \\ 7 & 5 & 6 & 5 \\ 8 & 6 & 10 & 9 \\ 7 & 5 & 9 & 10 \end{pmatrix} \begin{pmatrix} -7,2 \\ 14,6 \\ -2,5 \\ 3,1 \end{pmatrix} = \begin{pmatrix} 31,9 \\ 23,1 \\ 32,9 \\ 31,1 \end{pmatrix}$$



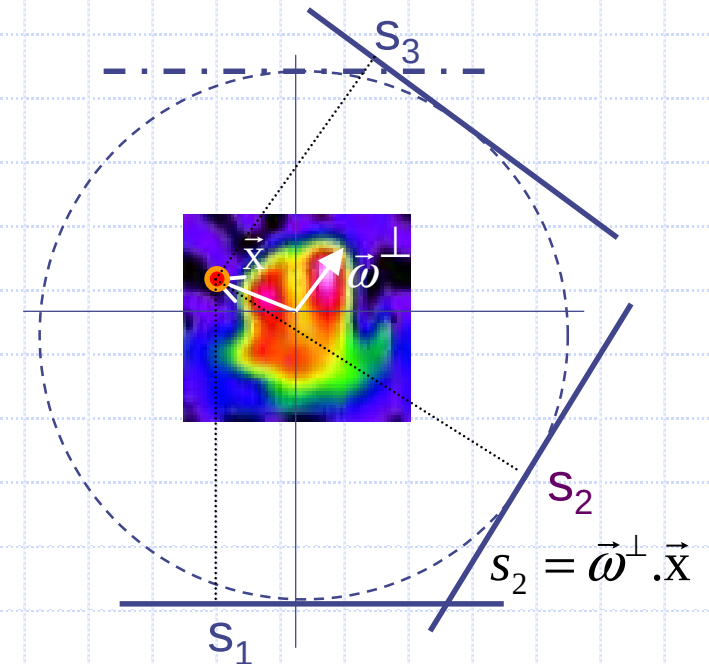
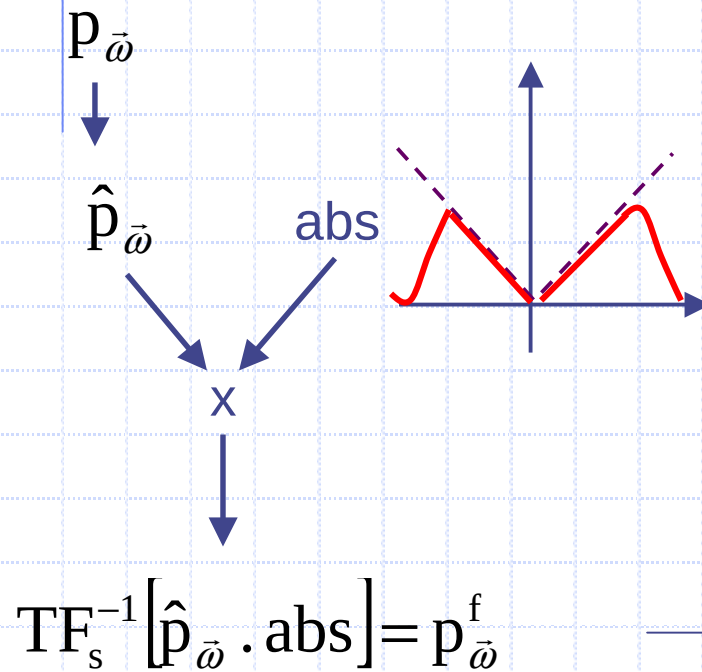
100 itérations de MLEM

Régularisation de la RPF



$$f(\vec{x}) = (R^* p^f)(\vec{x})$$

Projections sur 180°



Régularisation algébrique

Remplacer : $\bar{f} = \arg \min_{f \in C} \|\vec{p} - R\vec{f}\|^2$ par

$$\bar{f} = \arg \min_{f \in C} \left\{ \|\vec{p} - R\vec{f}\|^2 + \alpha \cdot \rho(\vec{f}) \right\}$$

Adéquation aux données
Surjectivité du problème inverse

Régularisation
injectivité

Exemple : régularisation de Tikhonov (cf. pseudo-inverse de Moore-Penrose)

$$\vec{f} = \arg \min_{\vec{f}} \left\{ \|\vec{p} - R\vec{f}\|^2 + \alpha \cdot \|\vec{f}\|^2 \right\} \Leftrightarrow (R^t R + \alpha I) f = R^t p$$

$$\vec{f} = (R^t R + \alpha I)^{-1} R^t p$$

Régularisation MAP-EM-OSL

Bayes : $P(\vec{f} / \vec{p}) = P(\vec{p}/\vec{f}) \cdot P(\vec{f})/P(\vec{p}) = P(\vec{p}/\vec{f}) \cdot P(\vec{f})$

$$\vec{f} = \arg \min_{\vec{f}} \left[-\log P(\vec{p}/\vec{f}) - \log P(\vec{f}) \right]$$

Adéquation aux données

régularisation

$$\log \prod_i \frac{e^{-\tilde{p}_i} \tilde{p}_i^{p_i}}{p_i!}$$

$$f_i^{n+1} = f_i^n \cdot \frac{1}{\sum_{l'=1}^P r_{l',i}} \sum_{l=1}^P r_{l,i} \frac{p_l}{\sum_{s=1}^N r_{l,s} f_s^n}$$

$$\text{Gibbs : } P(\vec{f}) = \frac{1}{K} e^{-\beta \cdot \sum_{i,j} w_{i,j} \cdot V(f_i - f_j)}$$

$$f_i^{n+1} = f_i^n \cdot \frac{1}{\sum_{l'=1}^P r_{l',i} + \beta \cdot \partial U} \cdot \sum_{l=1}^P r_{l,i} \frac{p_l}{\sum_{s=1}^N r_{l,s} f_s^n}$$

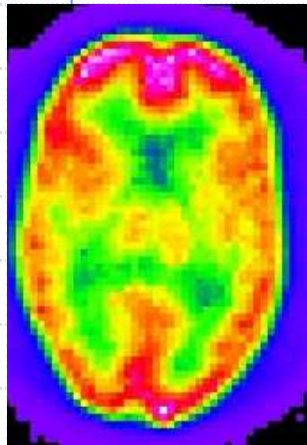
$$\partial U = \sum_{f_k \in V(f_i)} w_{i,k} \cdot \frac{\partial V}{\partial f} (f_i - f_k)$$

Dempster A et al. Maximum likelihood from incomplete data via the EM algorithm. J R Stat Soc 1977;39:1-38.

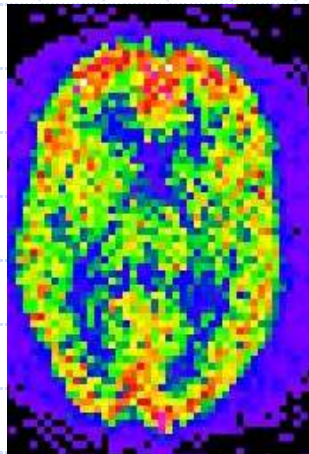
Hudson H et al. Accelerated image reconstruction using ordered subsets of projection data. IEEE TMI 1994;13:601-9.

Green PJ. Bayesian reconstructions from emission tomography data using a modified EM algorithm. IEEE Trans Med Imaging 1990;9:84-93

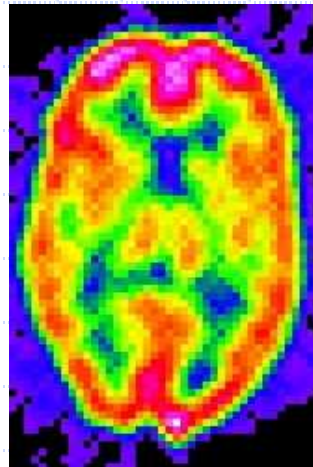
Comparaison MLEM-GC-FRECT



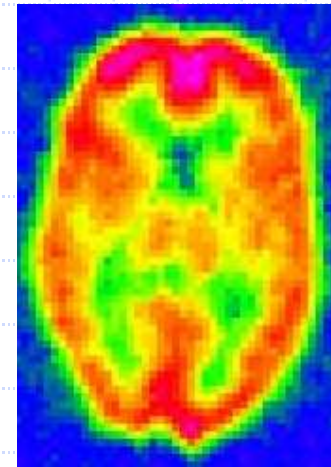
MLEM 6



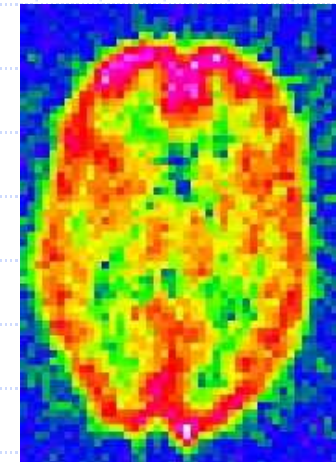
MLEM 200



FRECT 34 (CV)



GC 6



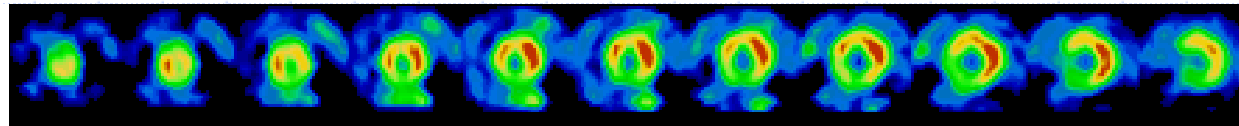
GC 16

$$E(f) = \left\| TF_s^{-1}(B.\hat{p}) - Rf \right\|^2 + \left\| (1 - B).\hat{f} \right\|^2$$

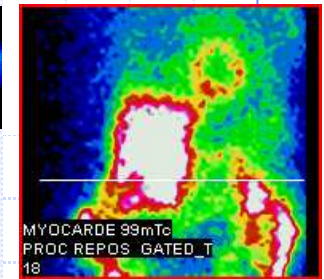
1. Problème médical à résoudre
2. Choix d'une modalité d'imagerie
3. Choix d'un traceur
4. Acquisition des images
5. Reconstruction tomographique
6. Segmentation, analyse et validation (myocarde)
7. Segmentation, analyse et validation (ventriculo)

Segmentation et Analyse

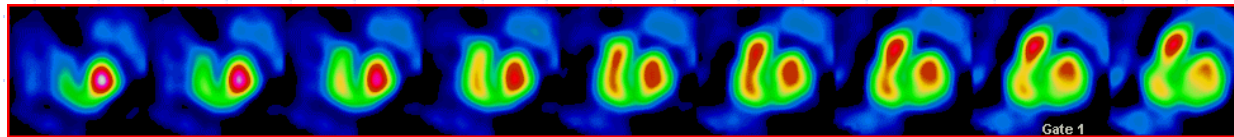
◆ Scintigraphie myocardique (MIBI ou MIBG)



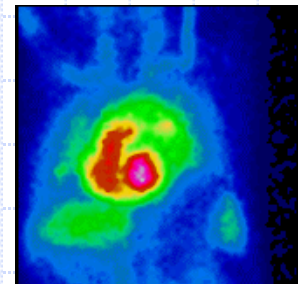
- ◆ Faible rapport signal/bruit
- ◆ Activités extra cardiaques
- ◆ Ventricule gauche $\approx$ ellipsoïde tronqué



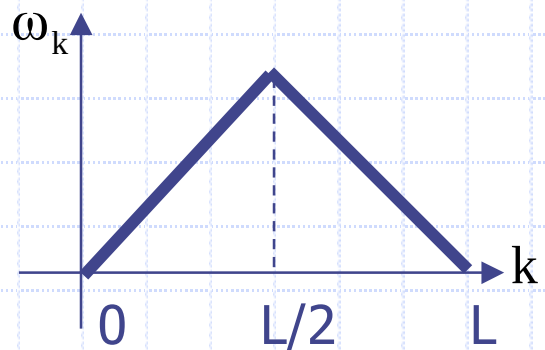
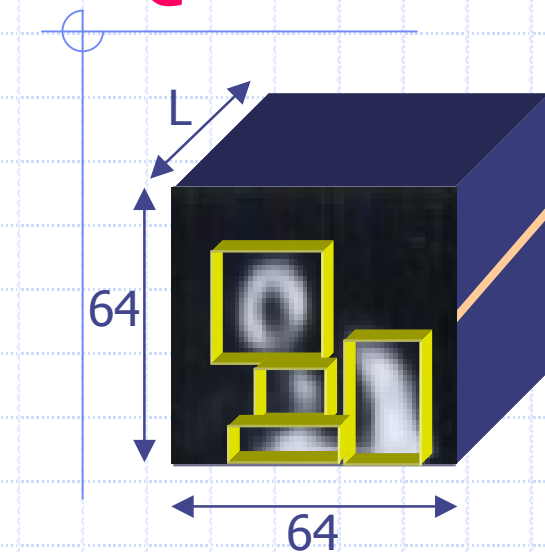
◆ Tomoventriculographie isotopique



- ◆ Meilleur rapport S/B, basse fréquence
- ◆ Volumes partiels
- ◆ Peu d'activité extra cardiaque
- ◆ Pas de modélisation simple (ventricule droit)



QPS® : MYOCARDE



Essai d'identification :
seuil → 1 amas cubique [50 mL, 1L[

S+5% ⇒ fragmentation

① Somme pondérée des coupes SA

$$S(i, j) = \sum_{k=1}^L \omega_k \cdot SA(i, j, k)$$

② ? Maxima locaux de $S(i, j)$

- supérieur seuil et au moins 4-connexes

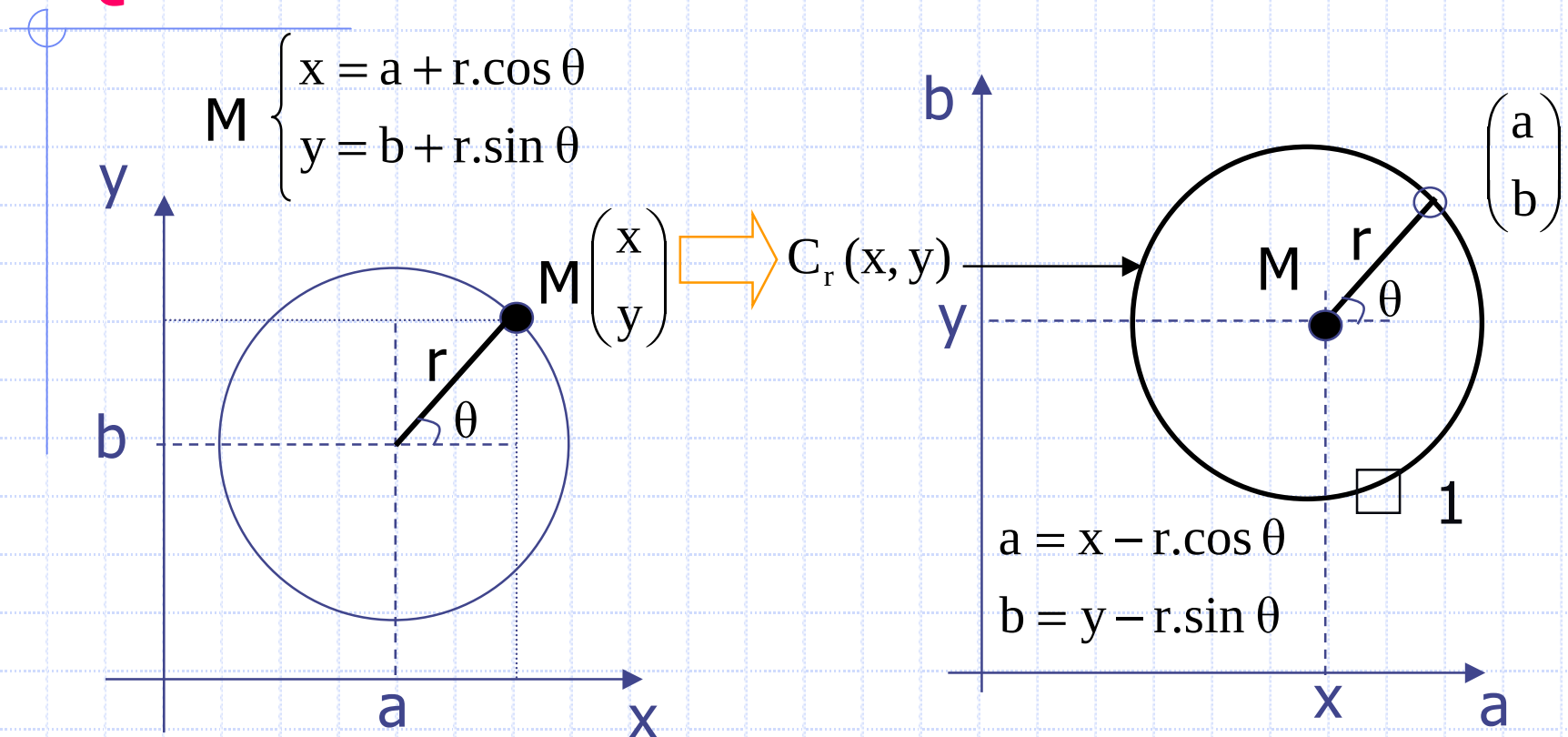
③ Transformée de Hough

- ? cercles dans un nuage de points

A Ezekiel et al. in Computers in cardiology. NY, IEEE Computer society. 1991:237-240

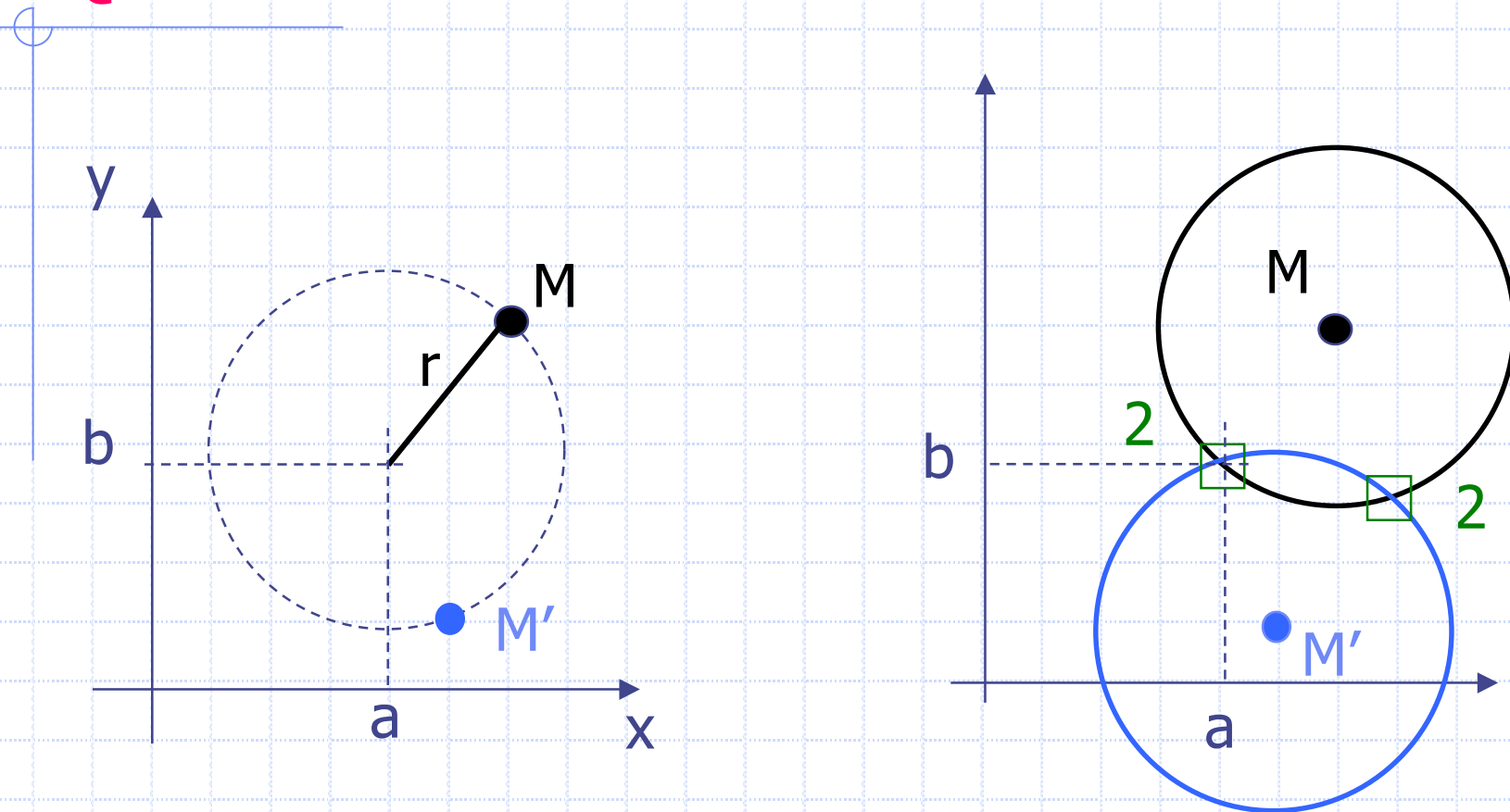
G. Germano et al. JNM 1995;36:1107-1114 et 1995;36:2138-2147

QPS[®] : TRANSFORMATION DE HOUGH



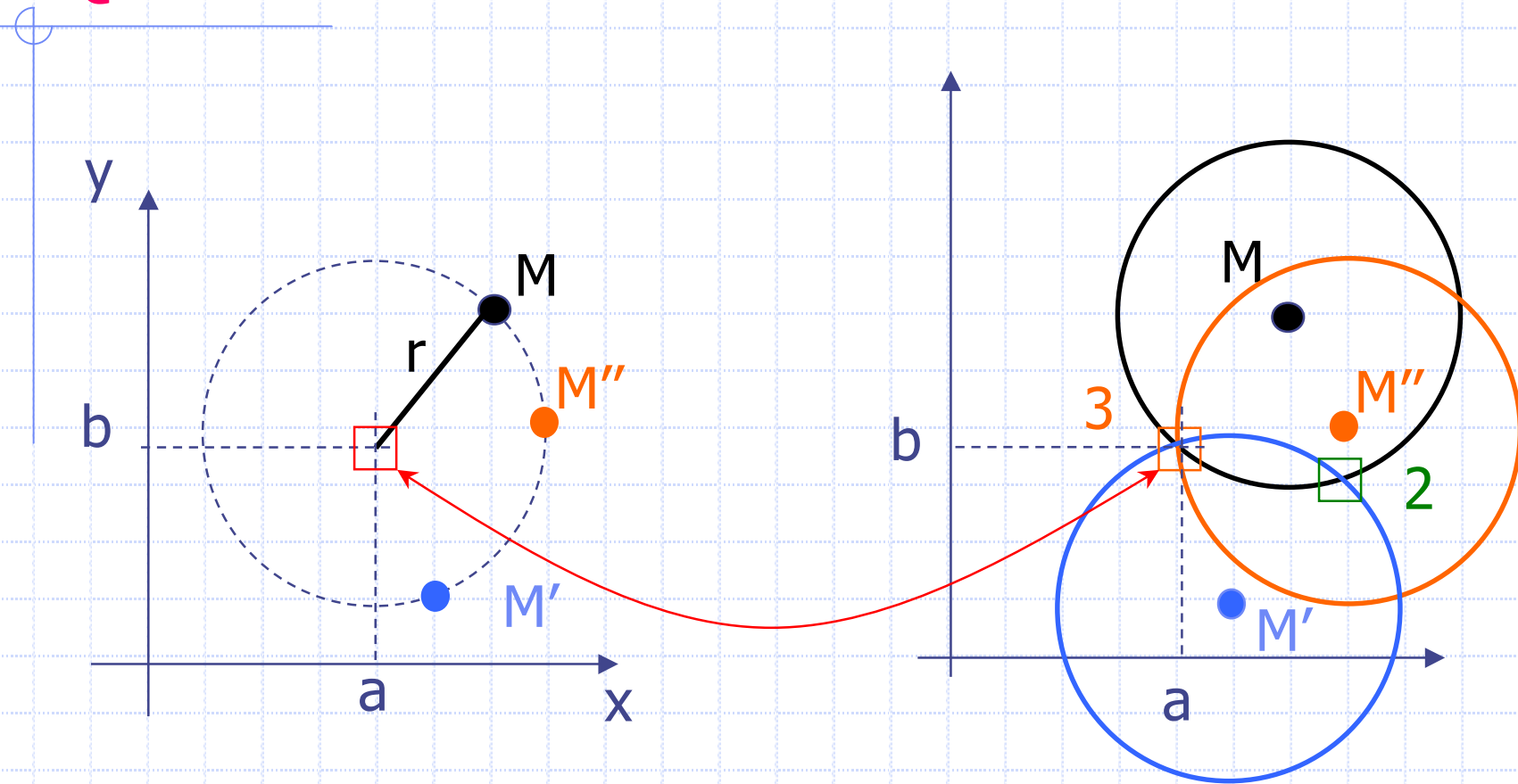
à tout **point** M du cercle(O,r) on associe un **cercle** (M,r)

QPS[®] : TRANSFORMATION DE HOUGH



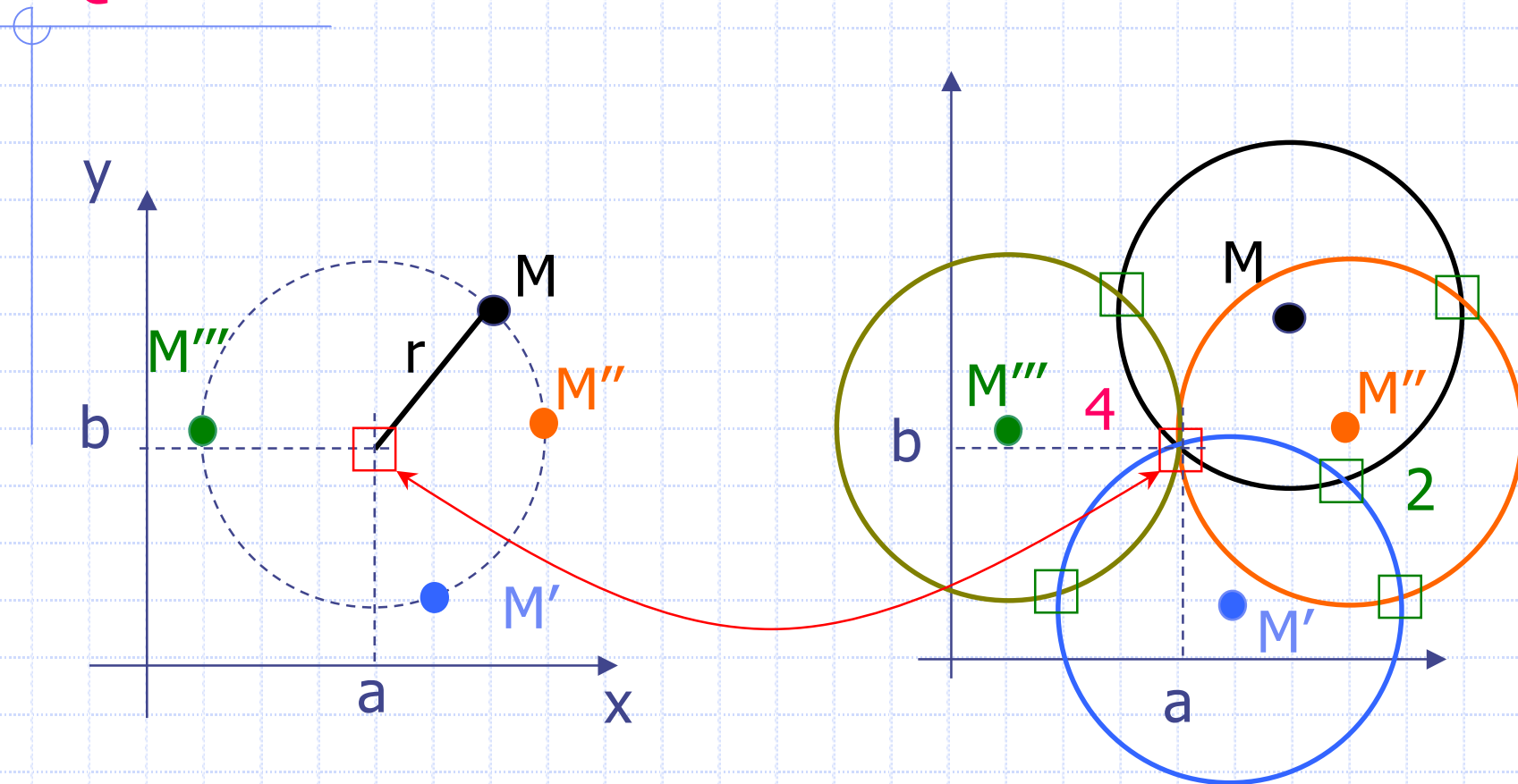
D. Ballard, C. Brown. Computer vision. Englewood Cliffs, NJ: Prentice-Hall; 1982.
<http://homepages.inf.ed.ac.uk/rbf/BOOKS/BANDB/bandb.htm>

QPS[®] : TRANSFORMATION DE HOUGH



D. Ballard, C. Brown. Computer vision. Englewood Cliffs, NJ: Prentice-Hall; 1982.
<http://homepages.inf.ed.ac.uk/rbf/BOOKS/BANDB/bandb.htm>

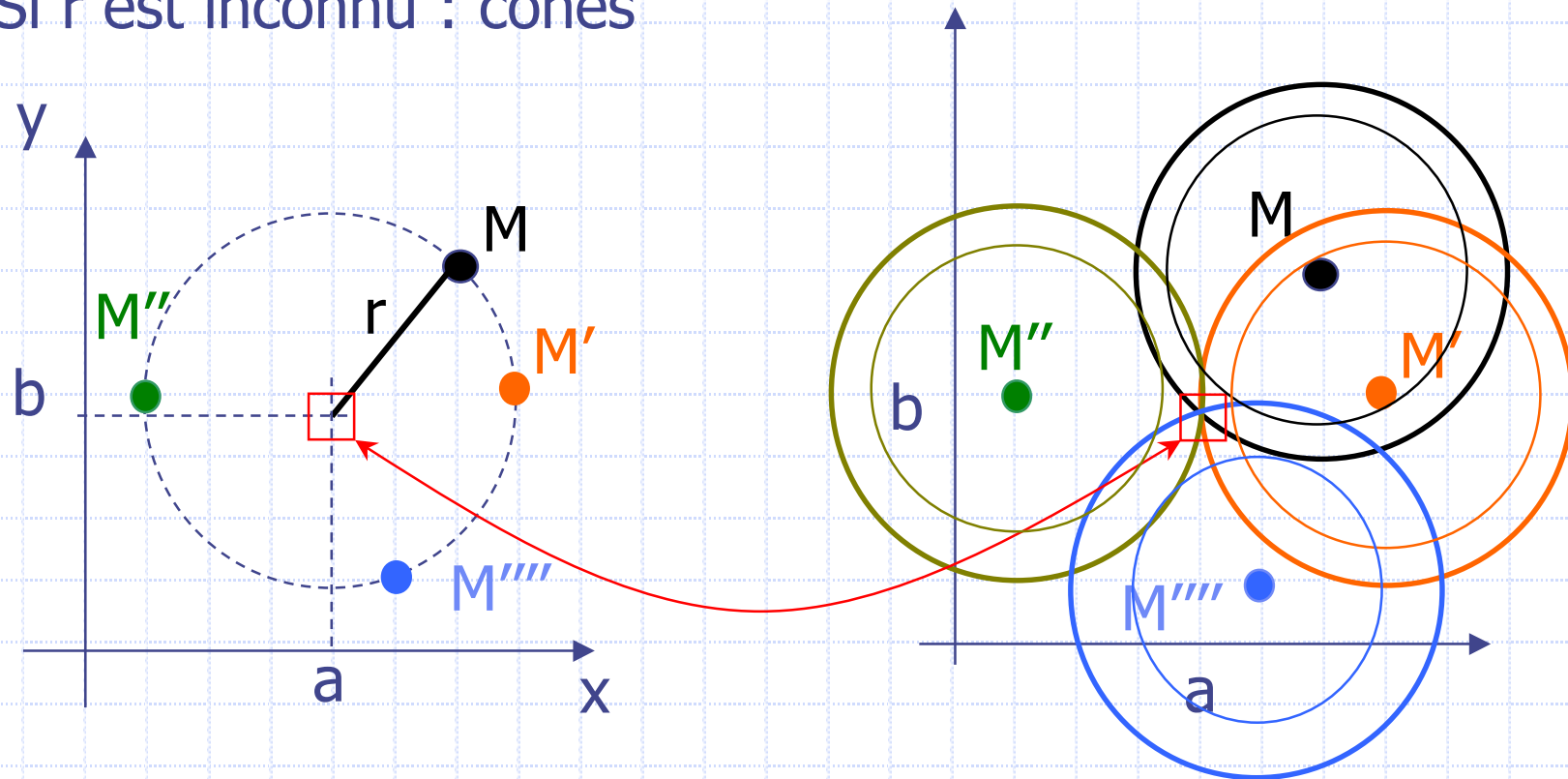
QPS[®] : TRANSFORMATION DE HOUGH



D. Ballard, C. Brown. Computer vision. Englewood Cliffs, NJ: Prentice-Hall; 1982.
<http://homepages.inf.ed.ac.uk/rbf/BOOKS/BANDB/bandb.htm>

QPS[®] : TRANSFORMATION DE HOUGH

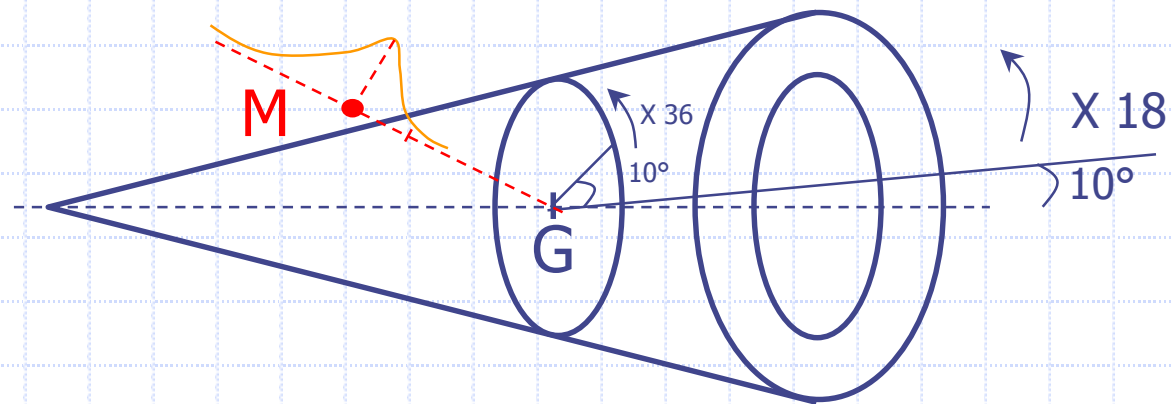
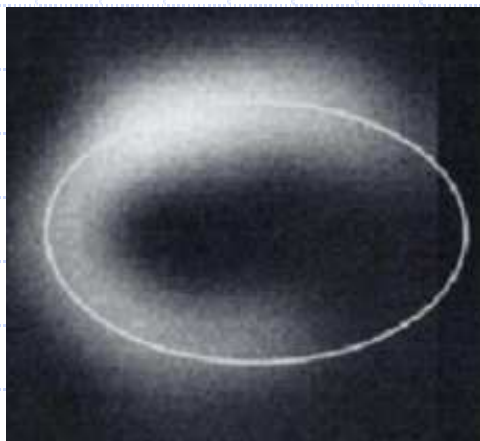
Si r est inconnu : cônes



? Extrema dans l'espace des paramètres 3D: (a, b, r)
Puis QPS choisit le cercle à plus fort comptage moyen
sur la circonférence par rapport au comptage au centre

QPS[®] : MYOCARDE

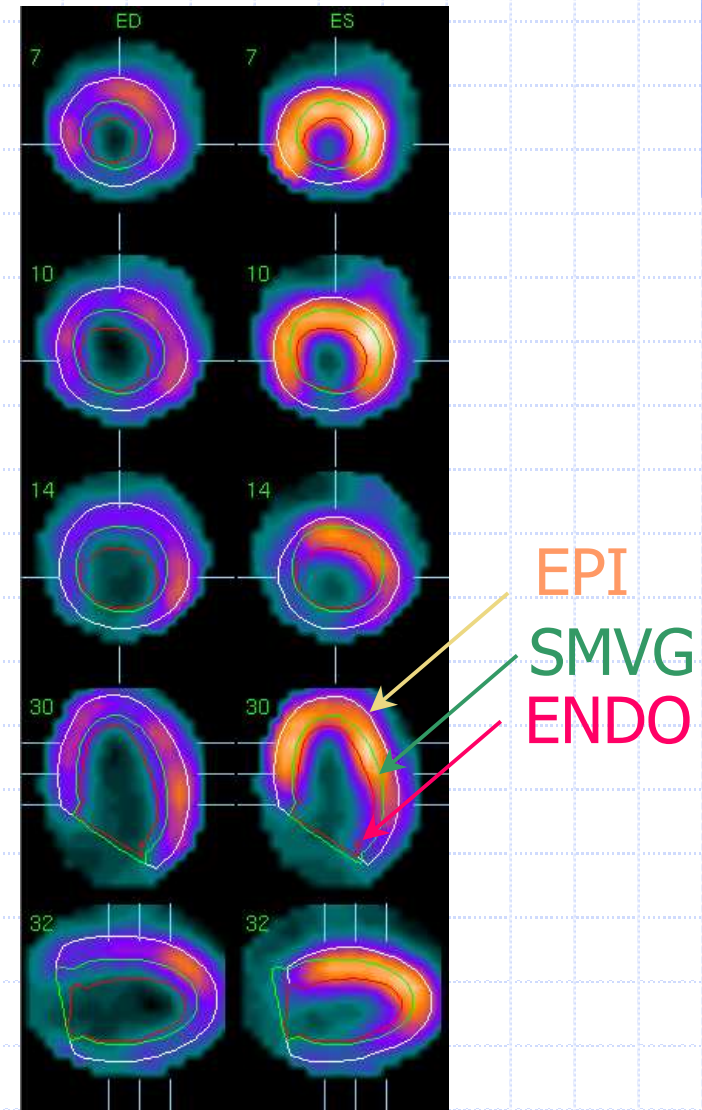
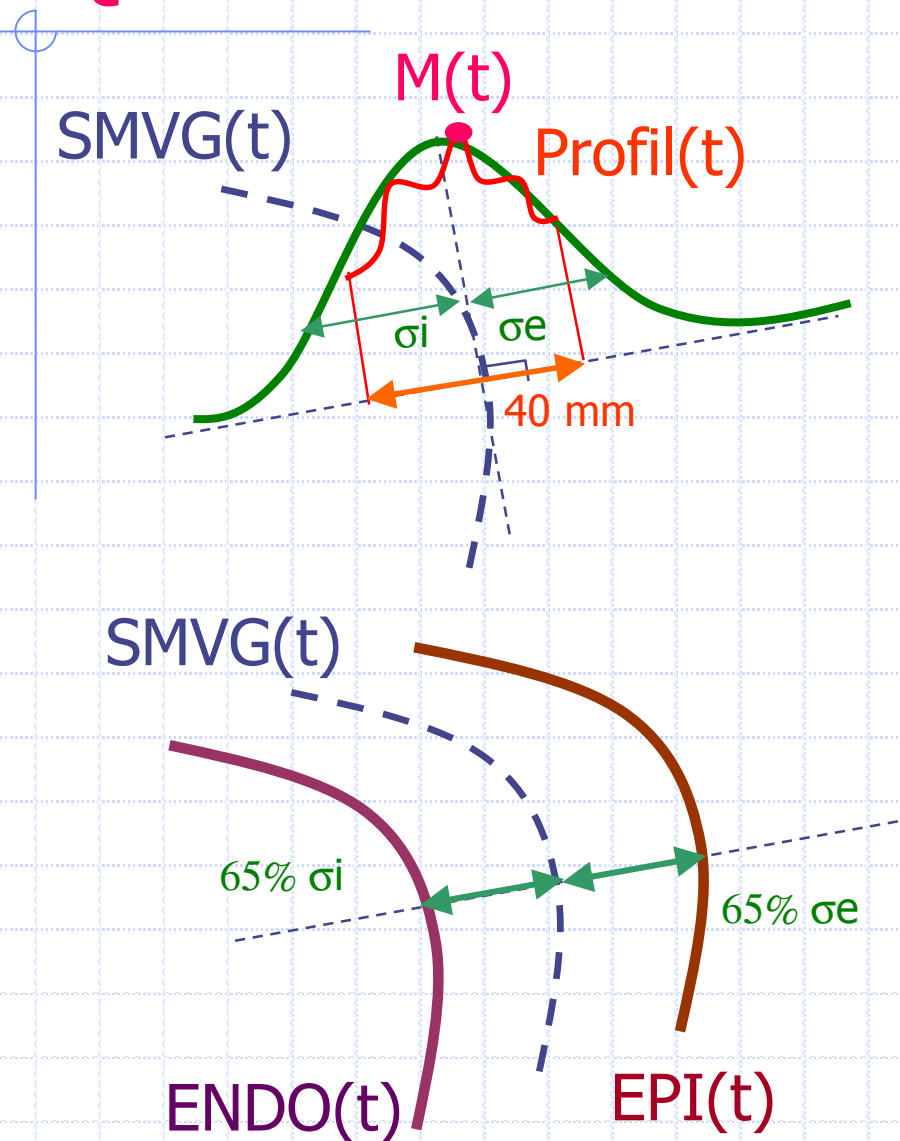
- ① G : Centre de Gravité du masque binaire
- ② Données = Volume SPECT x Masque
- ③ Extraction de 18x36 profils d'activité
- ④ Surface médio VG = maxima des profils M
- ⑤ Fit ellipse 3D → axe VG → G ← proj.(G)/axe



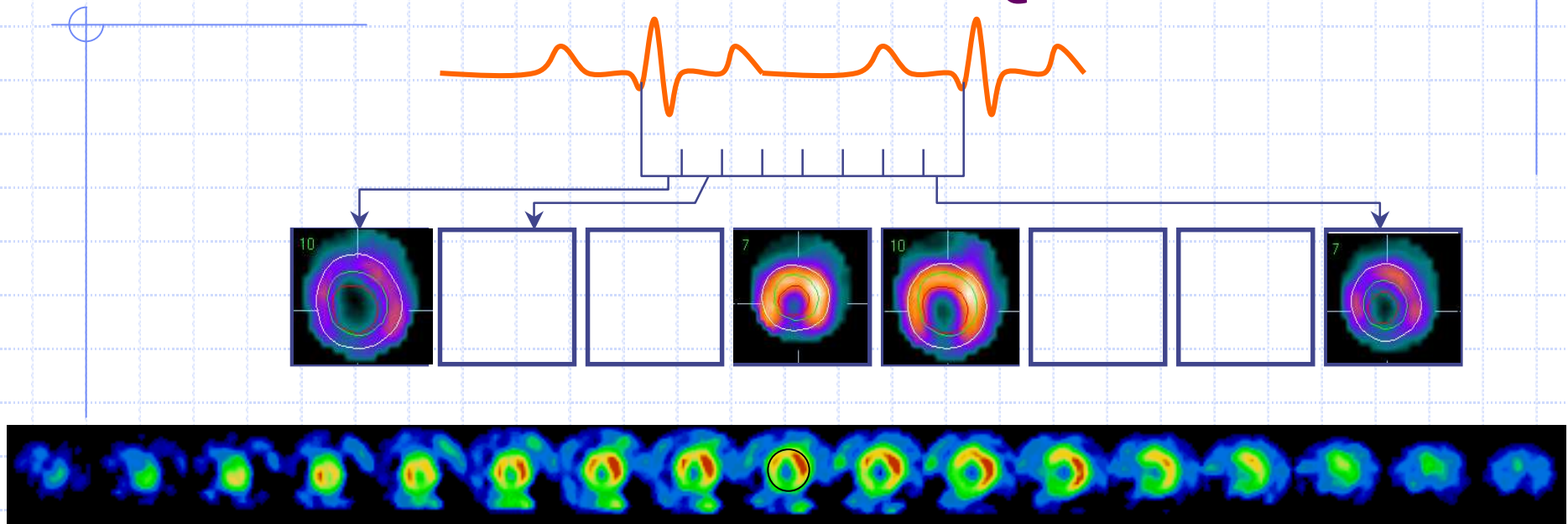
JC Cauvin et al. Eur J Nucl med 1992;19:1032-1037

G. Germano et al. JNM 1995;36:1107-14 ; 1995;36:2138-47 et JNM 2000;41:712-19

QPS[®] : SEGMENTATION DU MYOCARDE

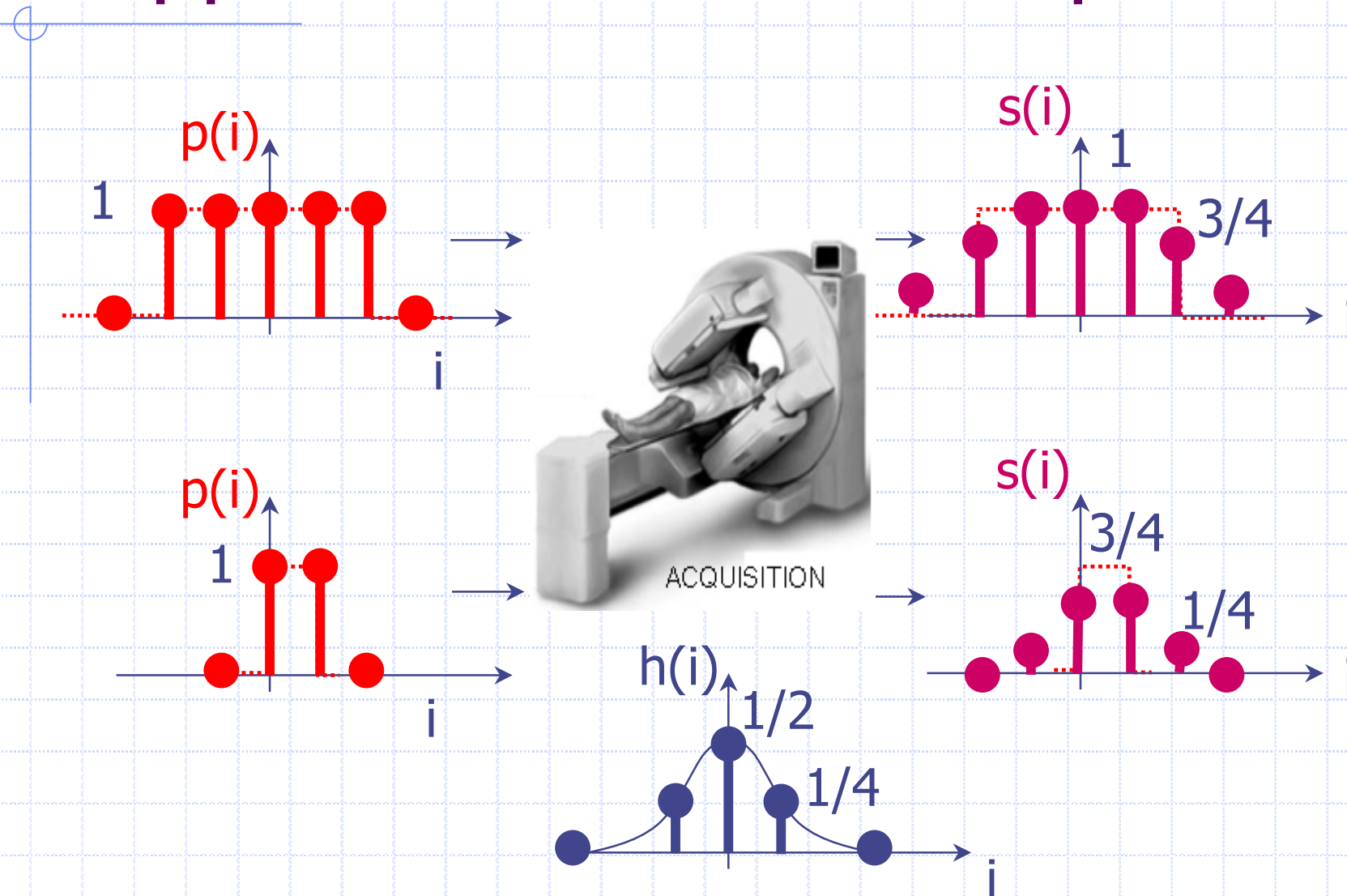


MOUVEMENT MYOCARDIQUE ?

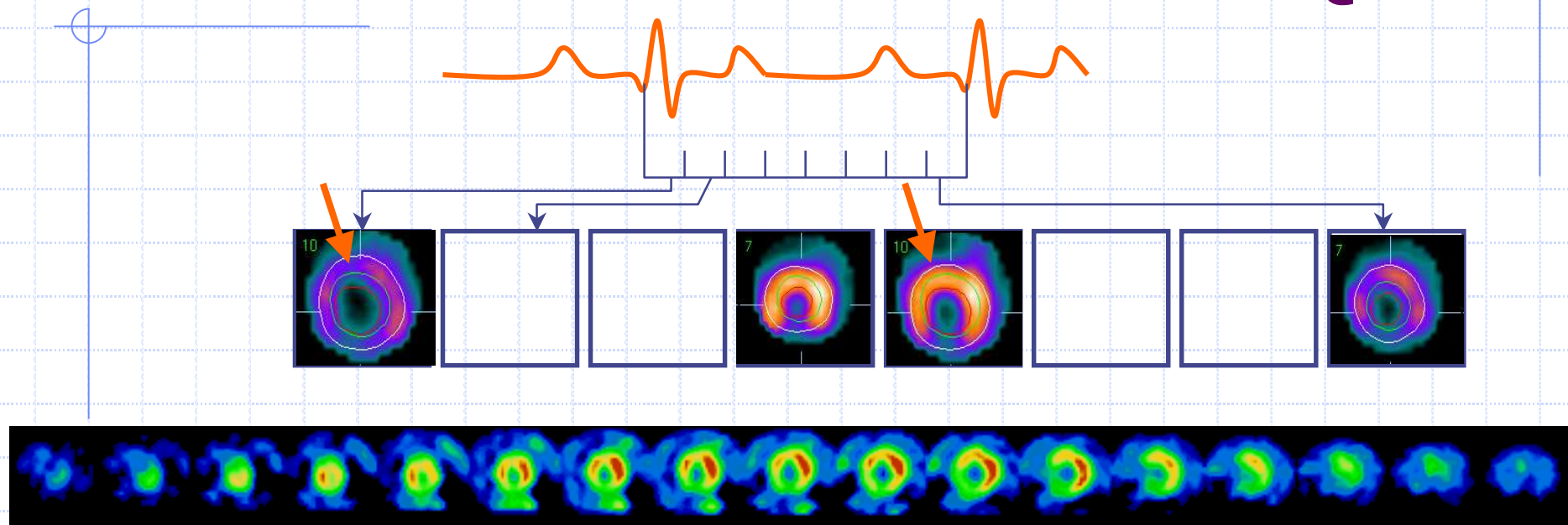


mais LMH ≈ 15 mm $\approx$ mouvement
qui est donc peu ou pas visible.

Rappel : Effet de volume partiel



EPAISSISSEMENT SYSTOLIQUE

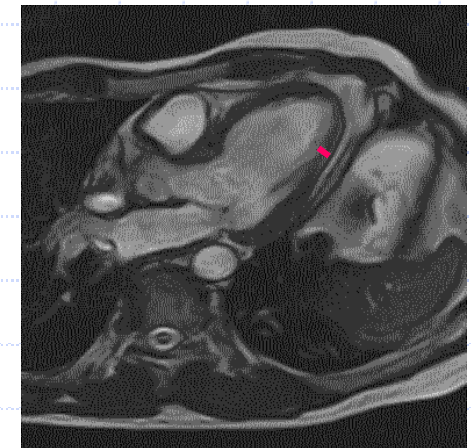


Epaisseur systolique ≈ 30 mm

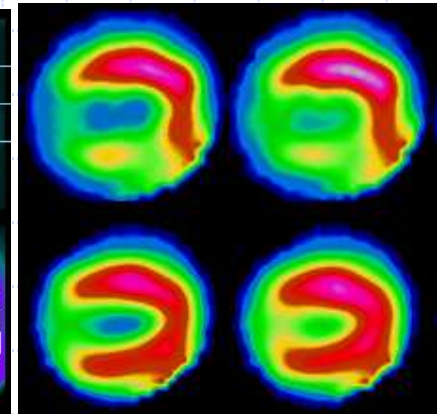
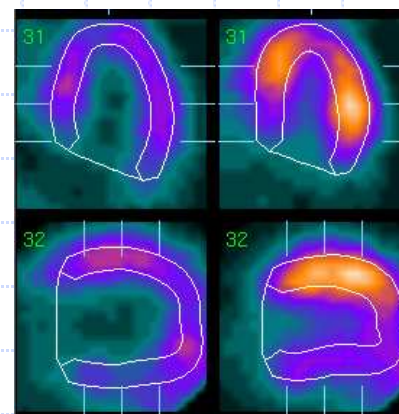
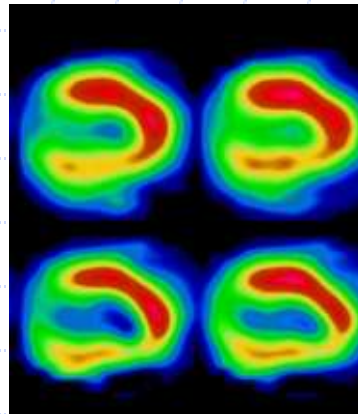
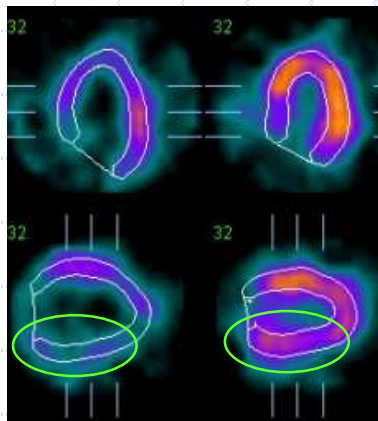
↪ 100 % de l'activité mesurée

Pour une épaisseur diastolique de 15 mm

↪ 75 % de l'activité mesurée



Applications cliniques



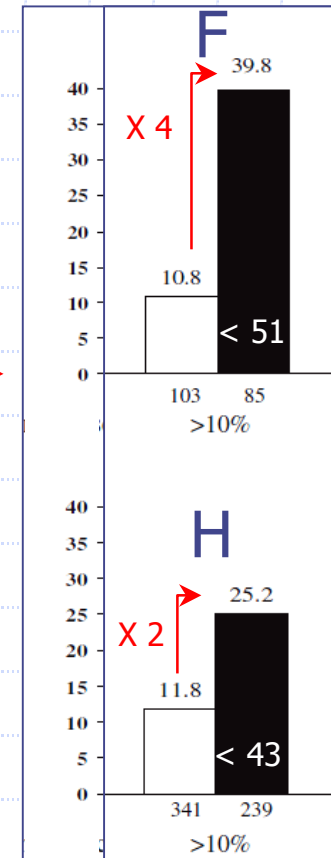
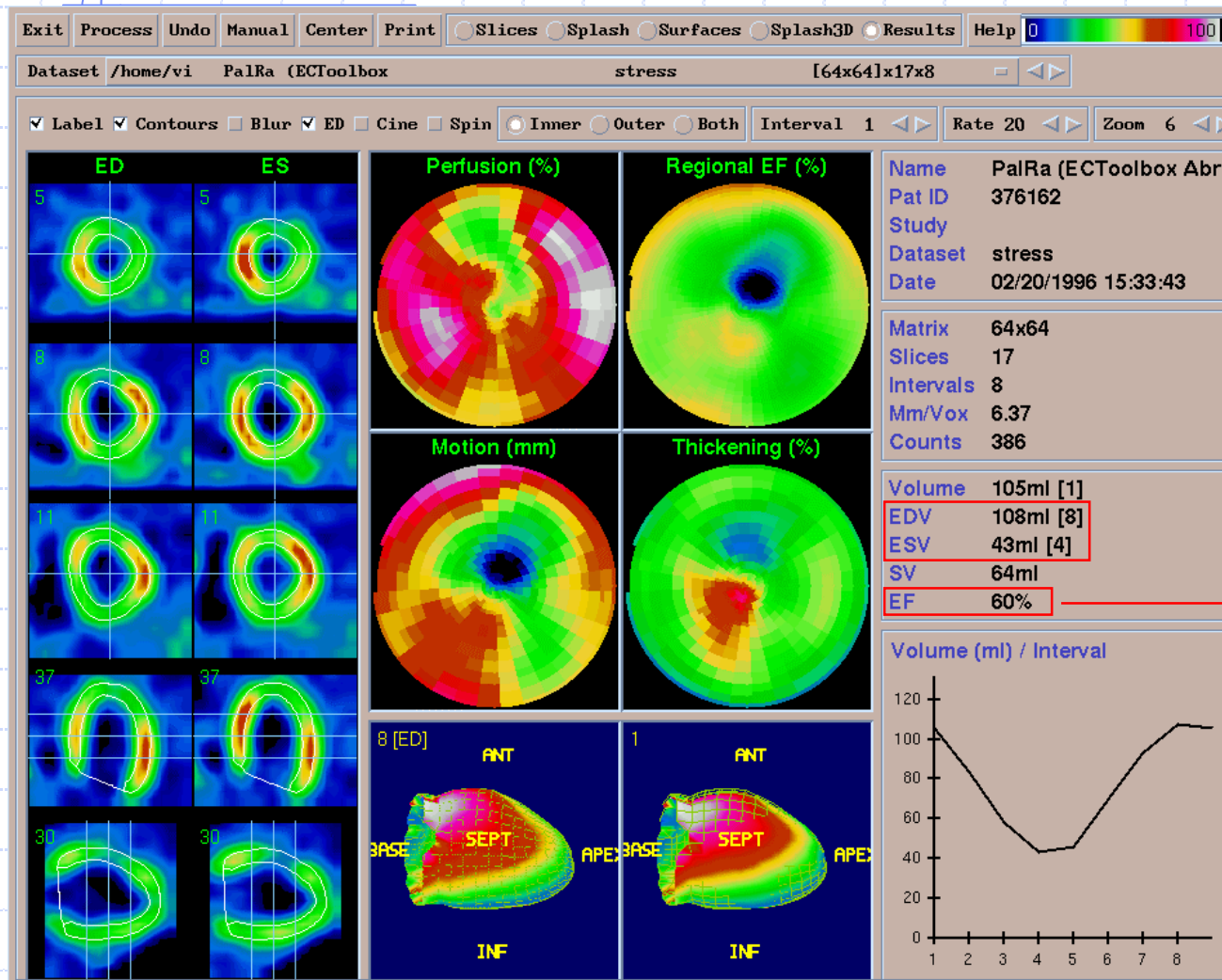
Examen normal

Se (%)	NC	88
p(T+/M)	ATT	86
Spé (%)	NC	50
p(T-/non M)	ATT	79



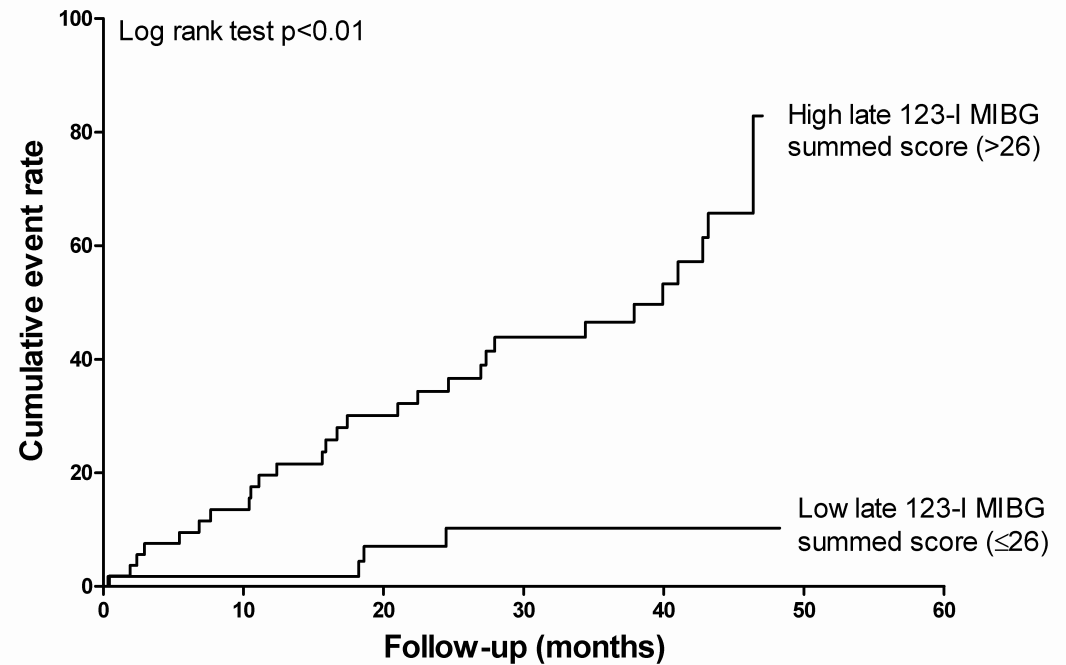
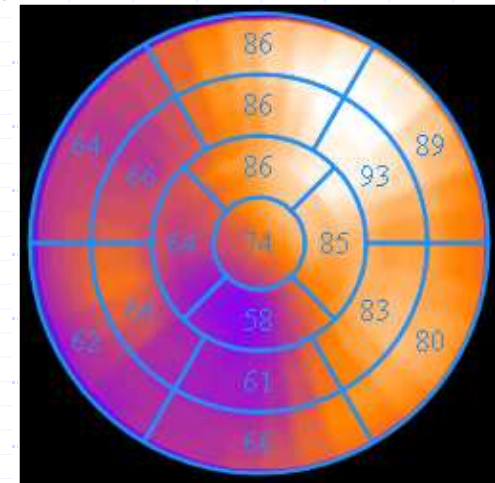
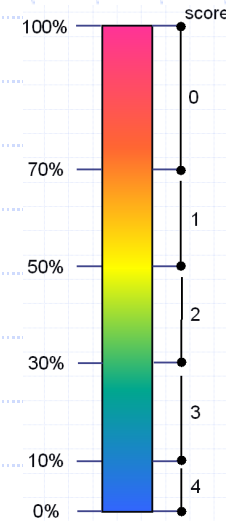
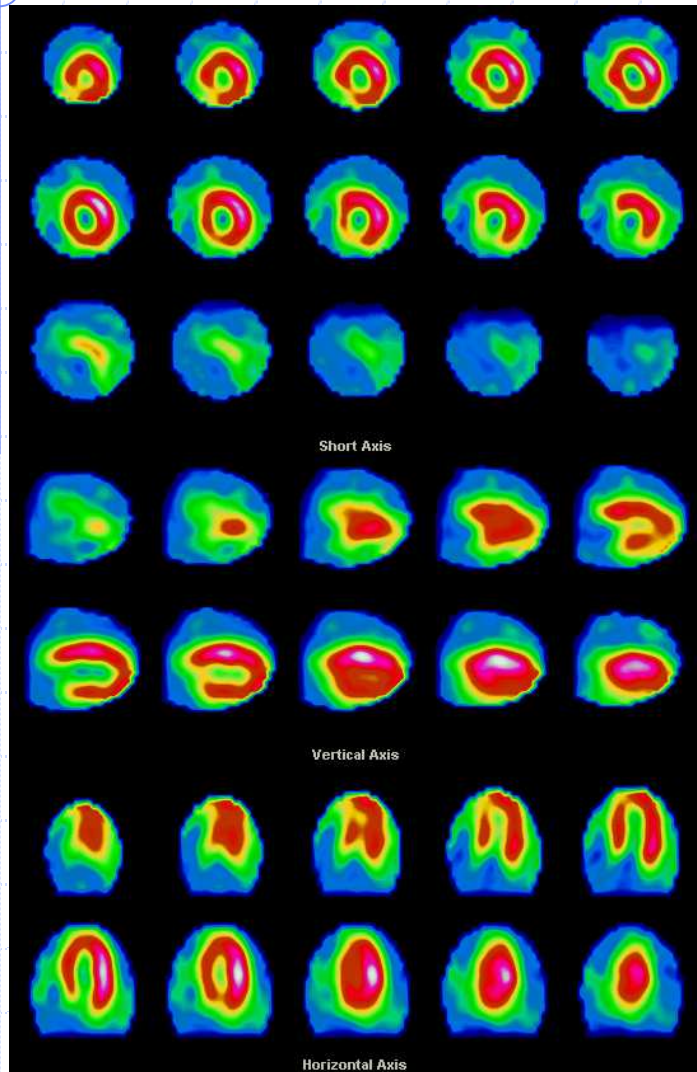
Sténose
de l'artère
circonflexe
> 80%

Applications cliniques



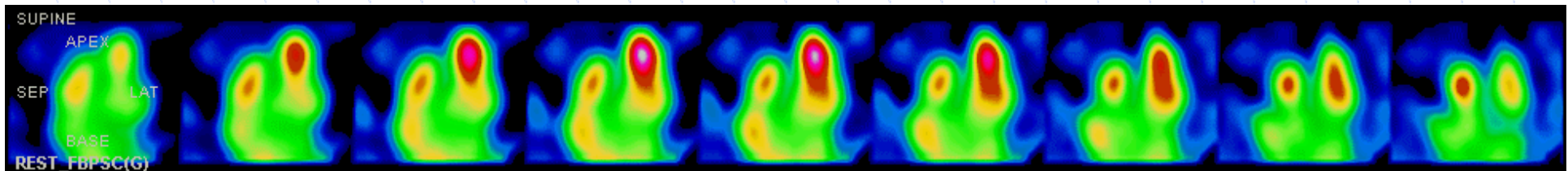
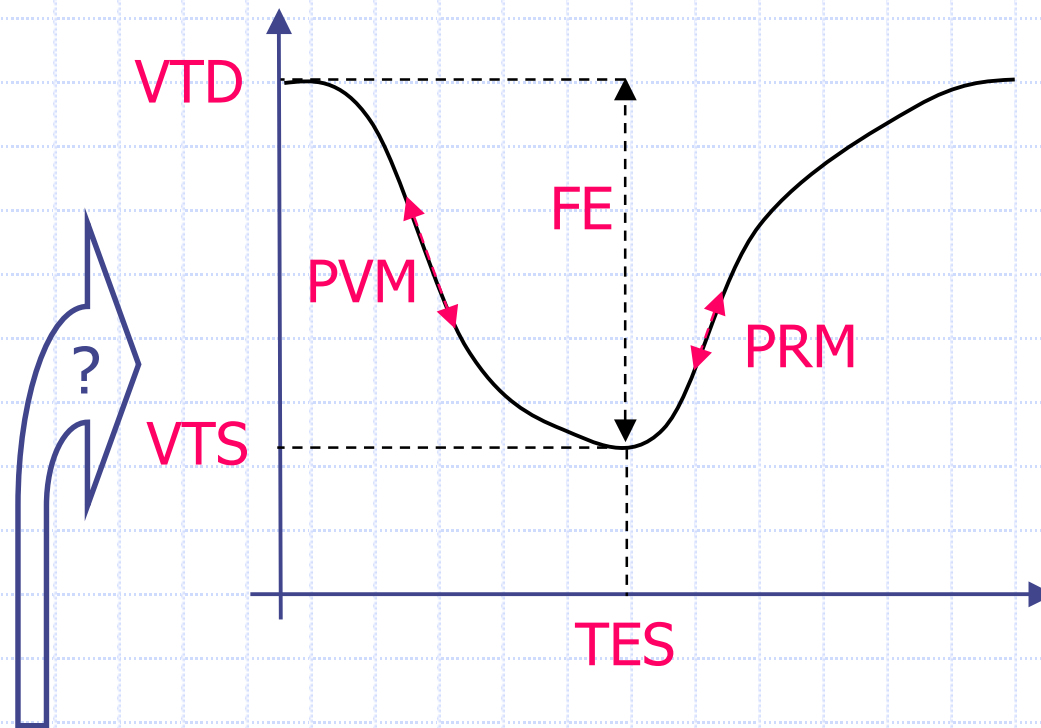
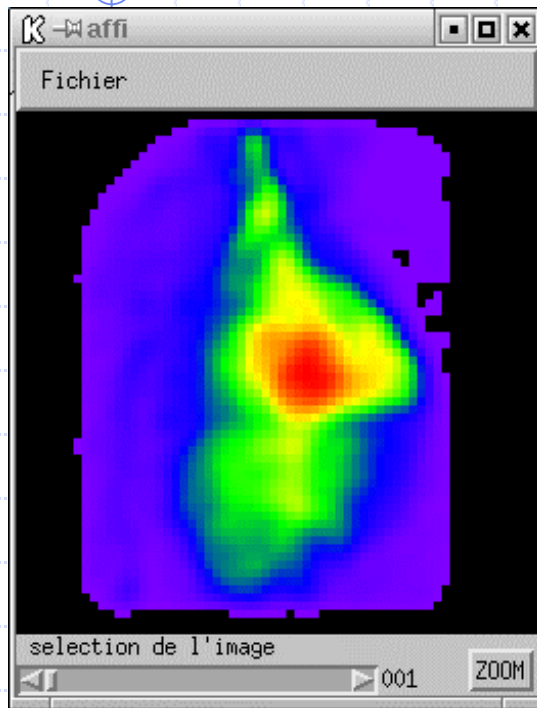
Evènements CV à 3 ans

MIBG



1. Problème médical à résoudre
2. Choix d'une modalité d'imagerie
3. Choix d'un traceur
4. Acquisition des images
5. Reconstruction tomographique
6. Segmentation, analyse et validation (myocarde)
7. Segmentation, analyse et validation (ventriculo)

TOMO-VENTRICULOGRAPHIE



Choix d'un algorithme de segmentation

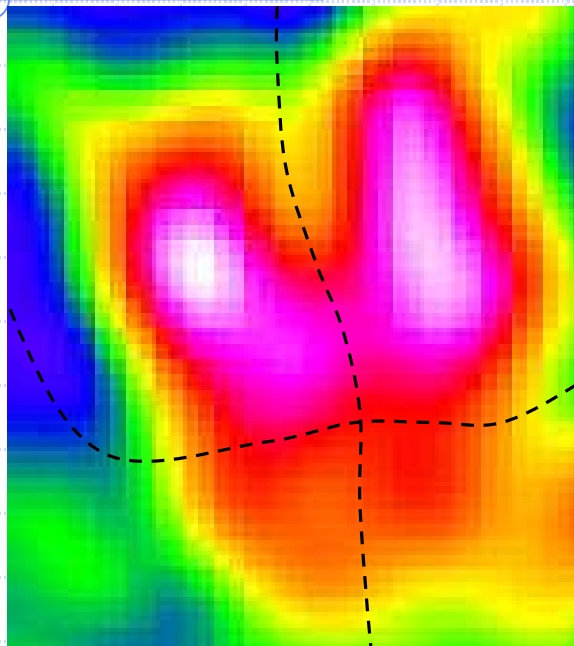
◆ Approche « frontières »

- ◆ Seuillages (simple, d'histogramme, hystérésis...)
- ◆ Dérivatives (gradient, Laplacien)
- ◆ Surfamiques (Hueckel)
- ◆ Morphologiques (gradient, ligne de partage des eaux)
- ◆ Réseau de neurones
- ◆ Champs de Markov
- ◆ Variationnelles

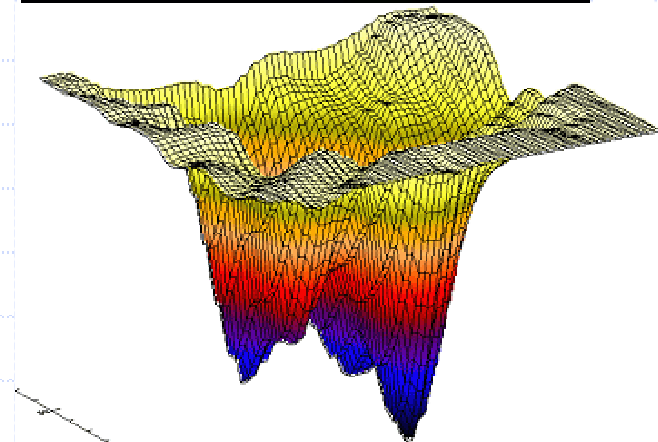
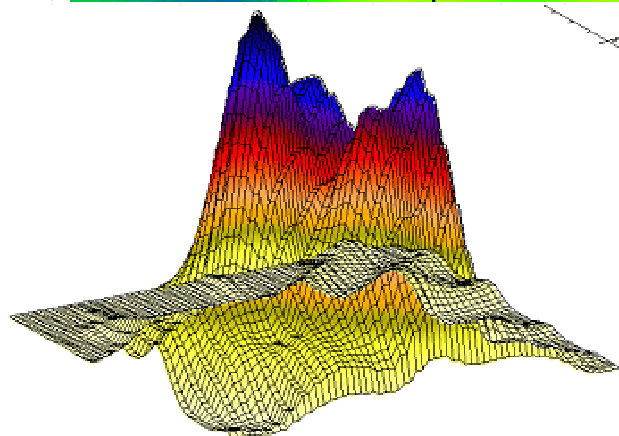
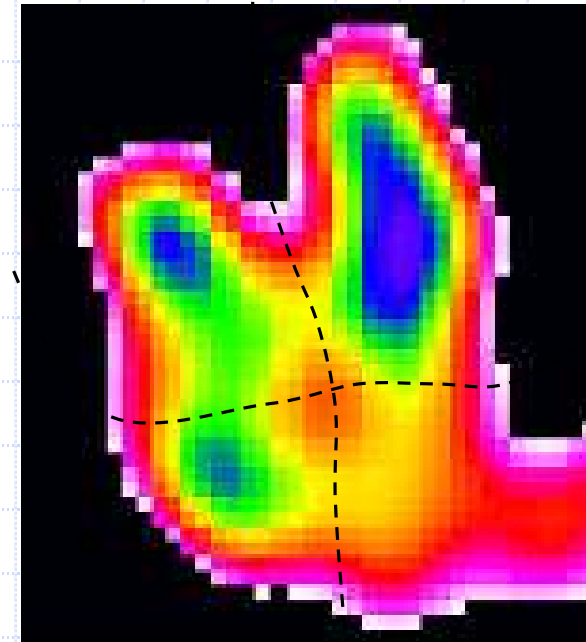
◆ Approche « régions »

- ◆ Techniques Markoviennes,
- ◆ Croissance de régions,
- ◆ Division et fusion de graphes,
- ◆ Etc.

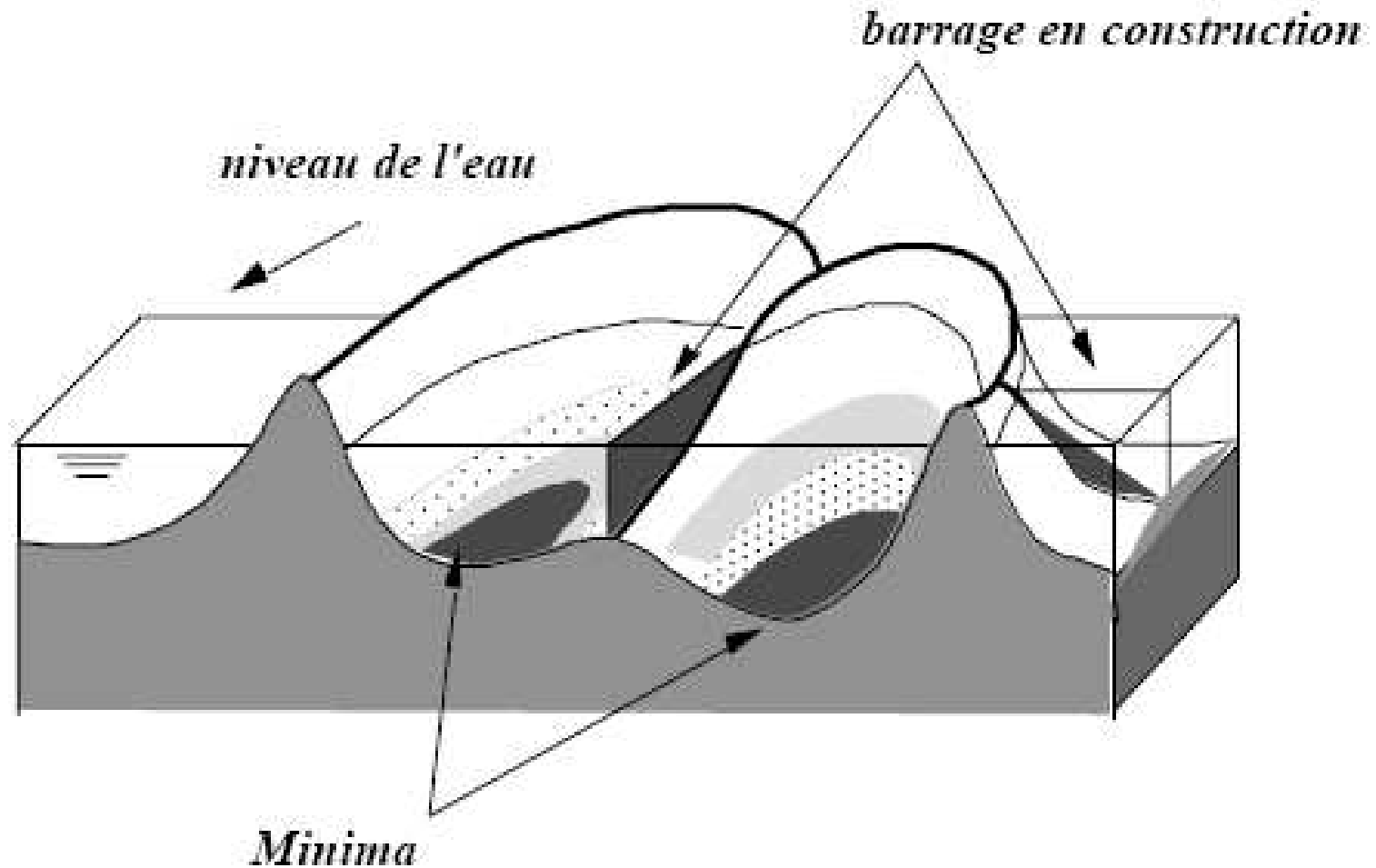
Segmentation



$$NG = NG_{\max} - NG$$

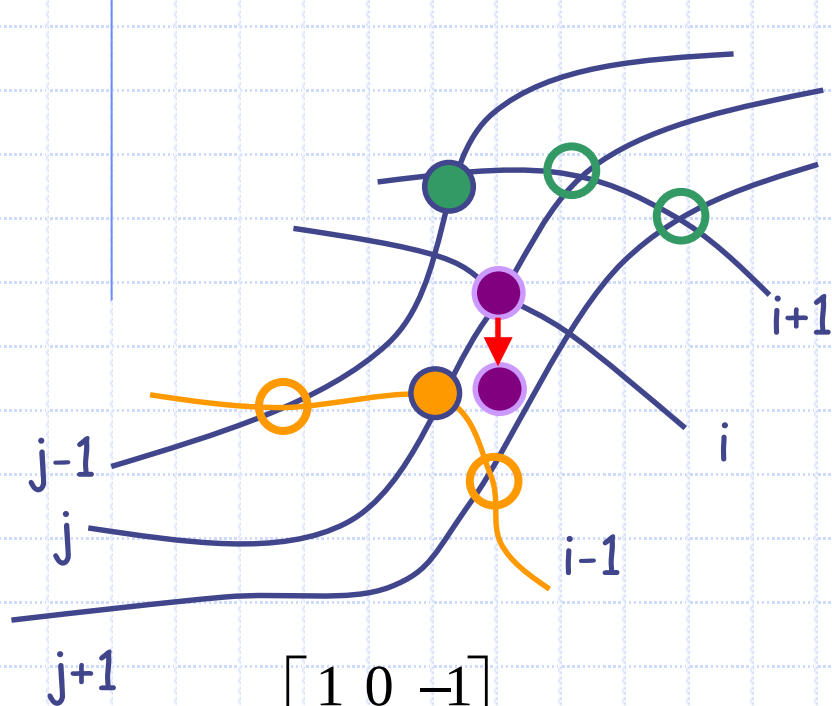


LPE par immersion

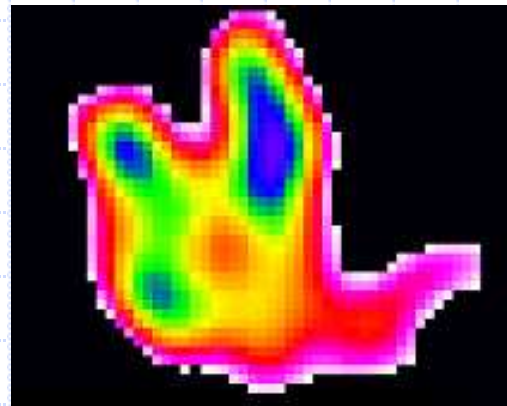


LPE par amincissement homotopique

si $f_{\max} < f(i, j) \leq f_{\min}$ alors $f(i, j) = f_{\max}$

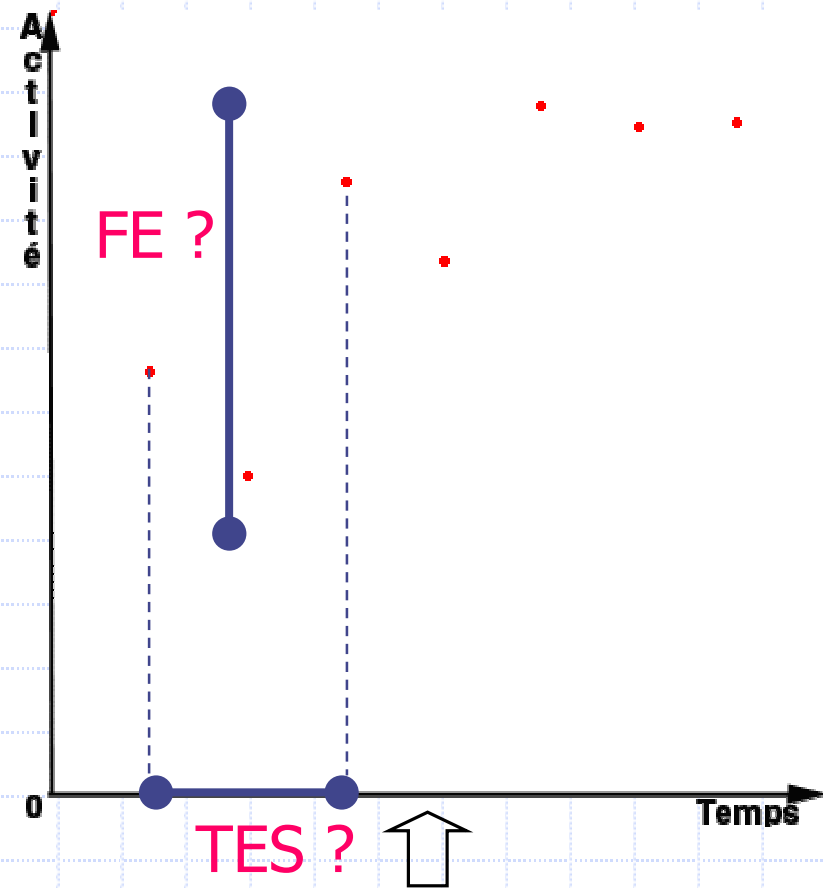
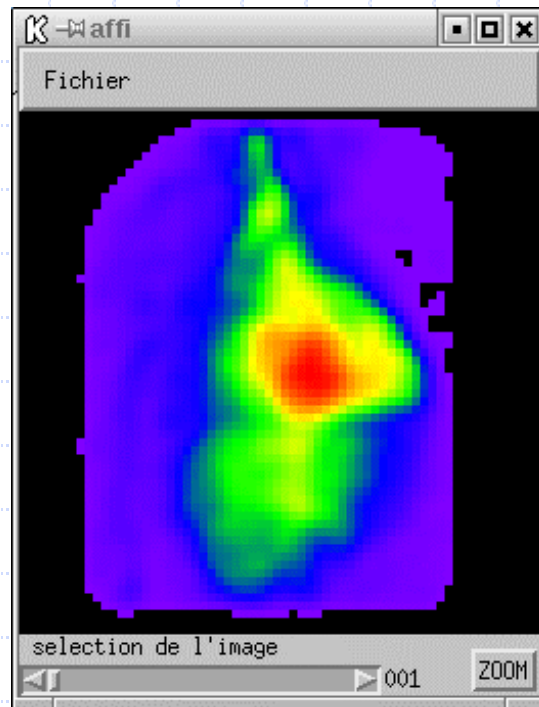


$$L = \begin{bmatrix} 1 & 0 & -1 \\ 1 & 0 & -1 \\ 1 & 0 & -1 \end{bmatrix}$$



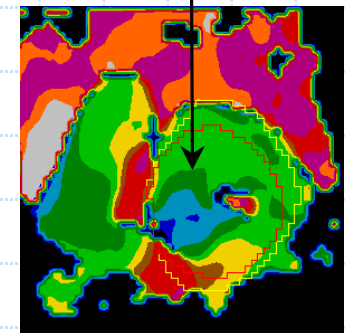
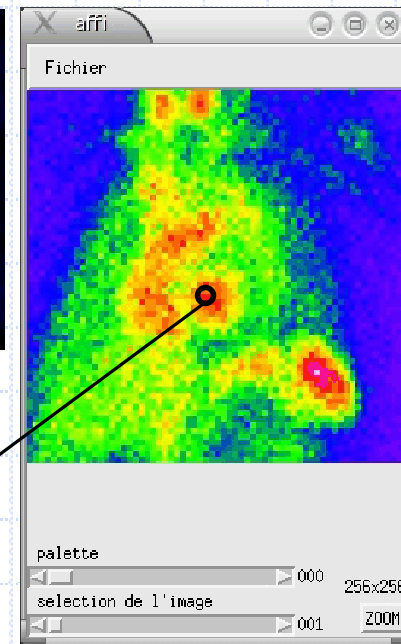
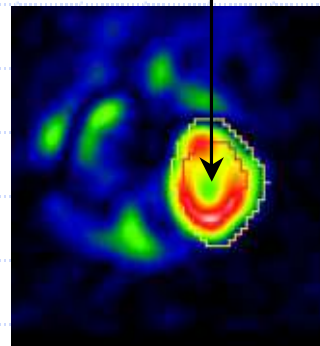
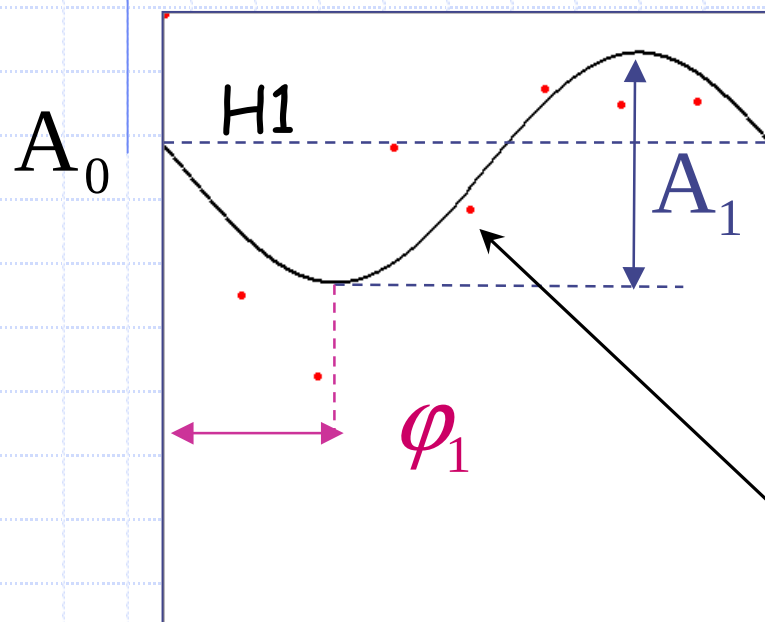
$$L' = \begin{bmatrix} 0 & 1 & 0 \\ 1 & -1 & 1 \\ 0 & 0 & 0 \end{bmatrix}$$

Extraction



Ajustement mono- harmonique

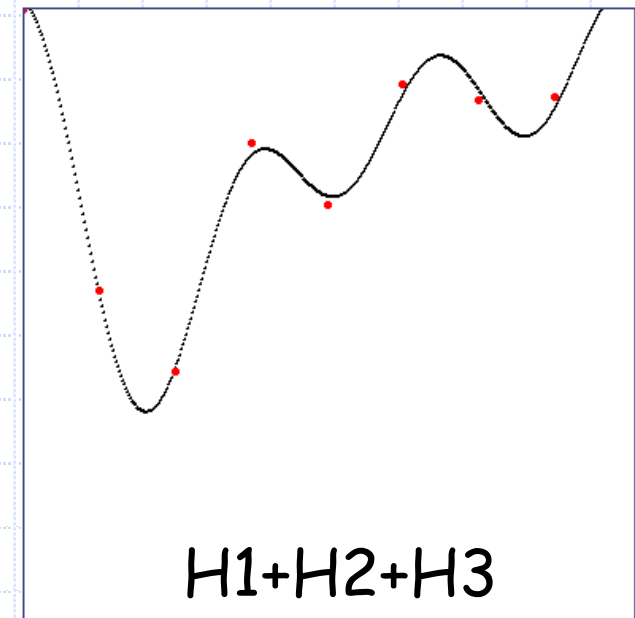
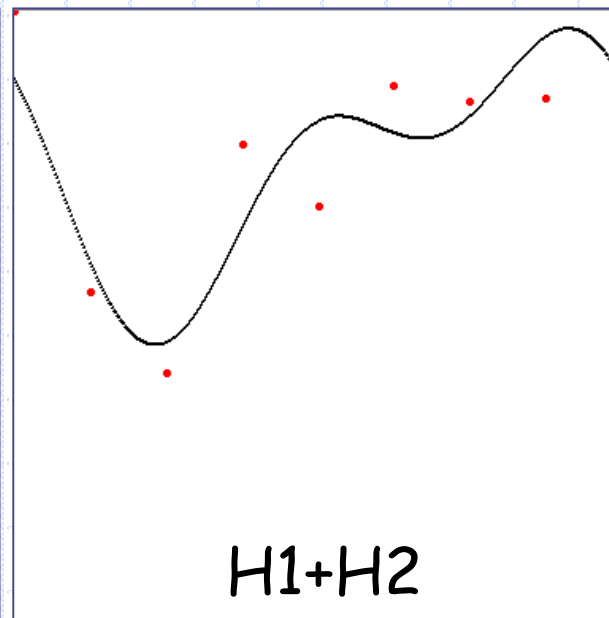
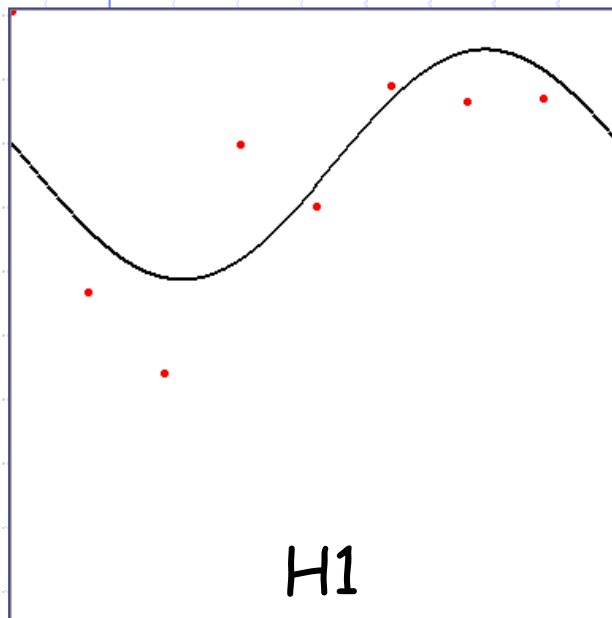
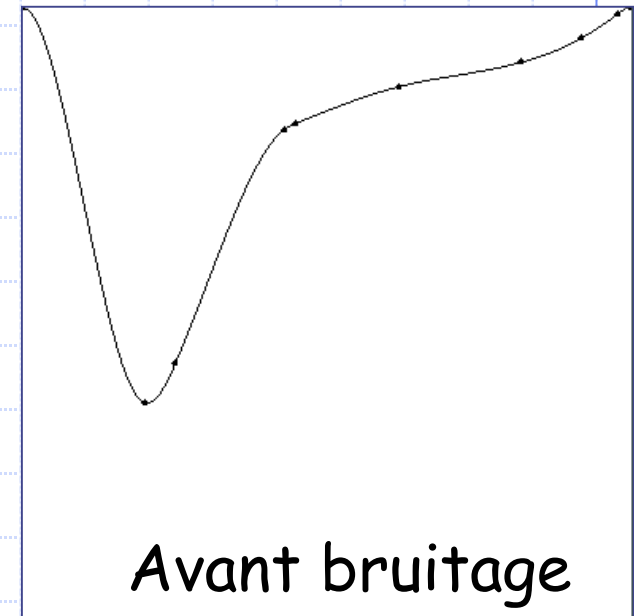
$$s(t) = \frac{\hat{s}(0)}{N} + \frac{\hat{s}(1)}{N} e^{j.(\omega_0)t} = A_0 + \underbrace{A_1 e^{j.(\omega_0 t + \phi_1)}}_{\text{fit cosinus}}$$



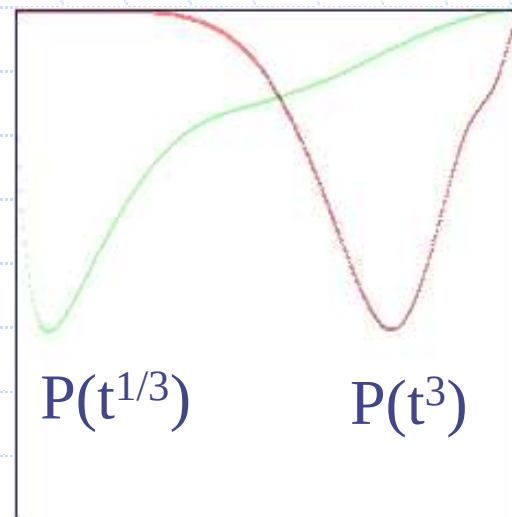
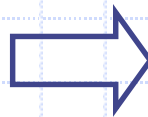
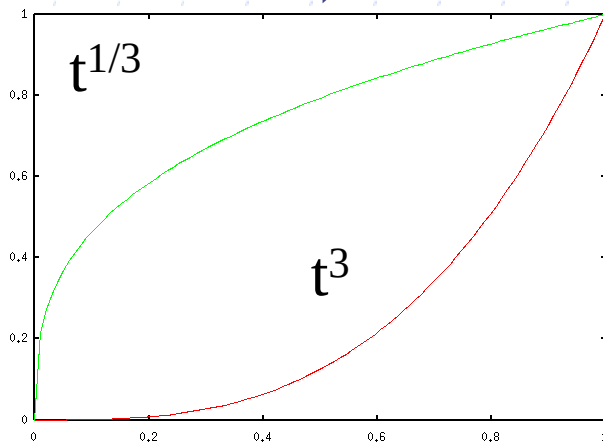
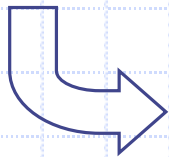
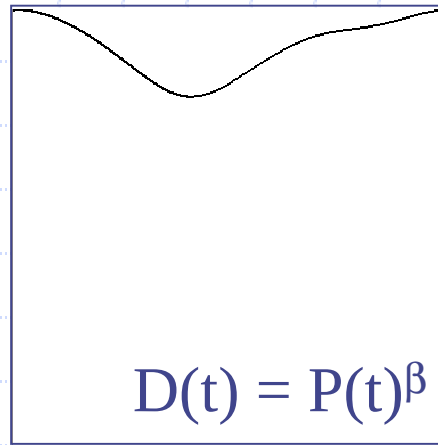
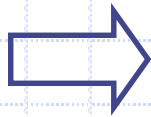
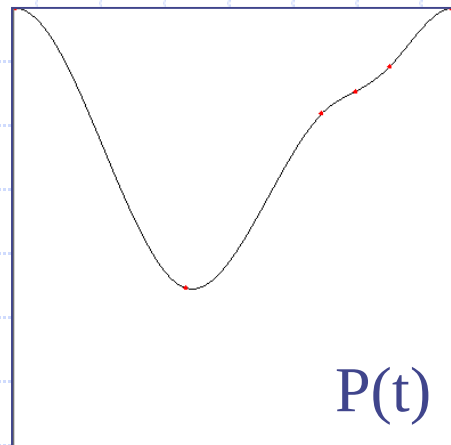
$$s(t) = \frac{1}{N} \sum_{k=0}^{N-1} \hat{s}(k).e^{j.(k.\omega_0)t}$$

Ajustement multi-harmonique

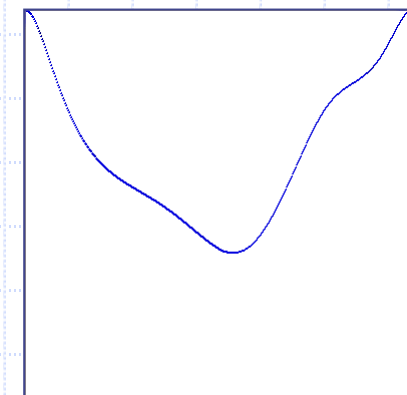
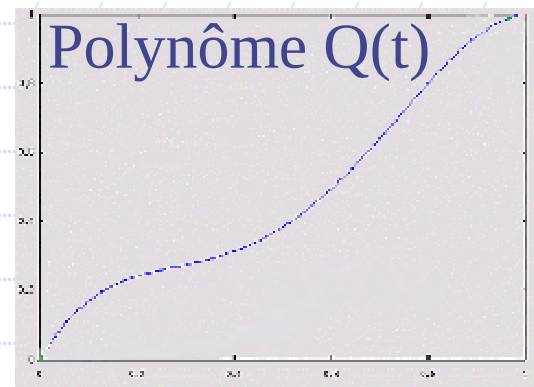
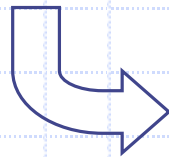
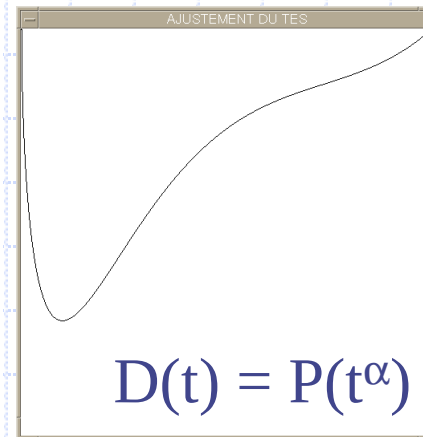
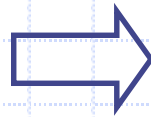
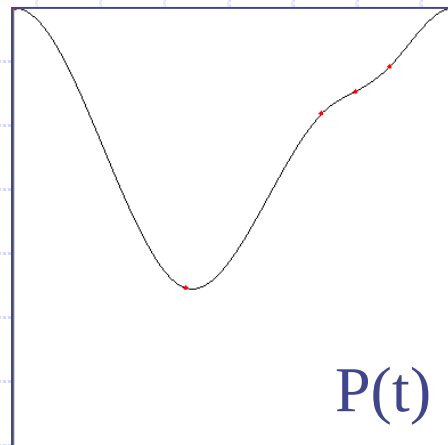
$$s(t) = \frac{1}{N} \sum_{k=0}^{H-1} \hat{s}(k) \cdot e^{j \cdot (k \omega_0) t}$$



Déformation d'une CTA de référence

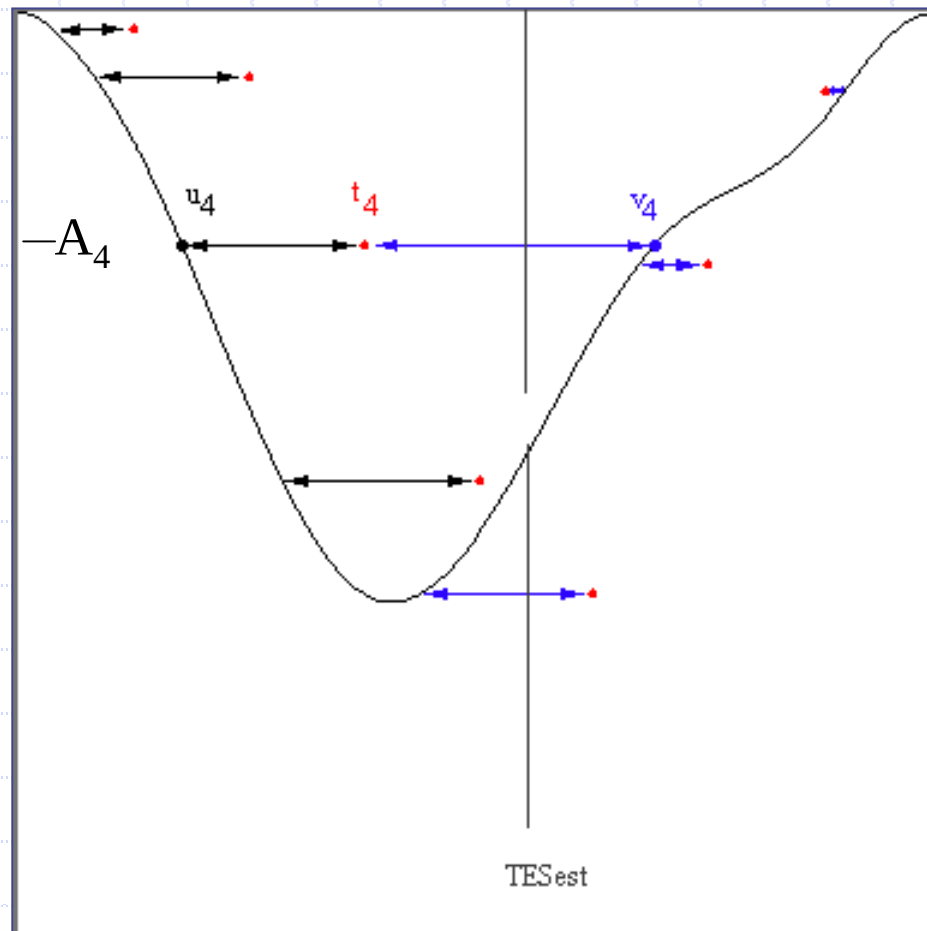


Déformation d'une CTA de référence



$$D(t) = P[Q(t)]$$

Détermination de $Q(t)$

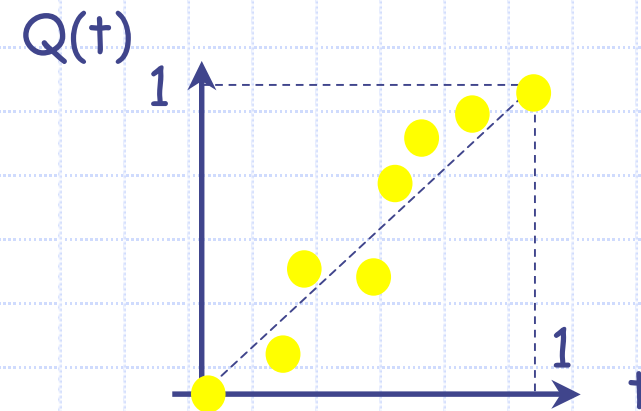


Acquisition bruitée (t_4, A_4)

$$A_4 = P(u_4)$$

$$A_4 = D(t_4) = P[Q(t_4)]$$

$$Q(t_4) = u_4$$

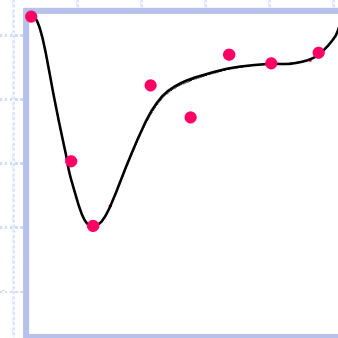
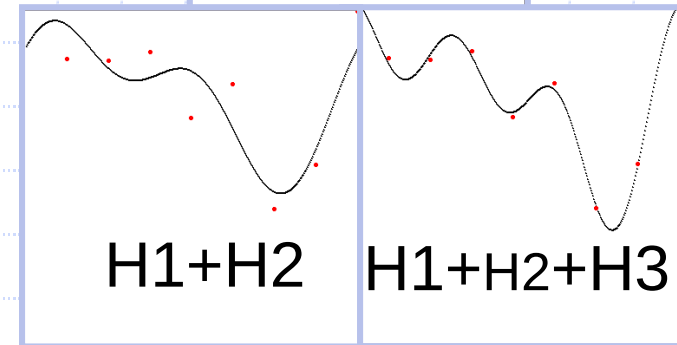
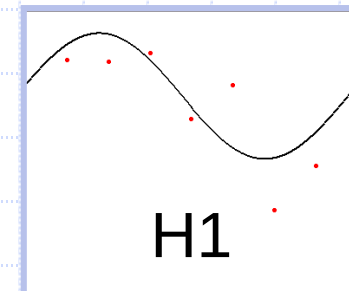
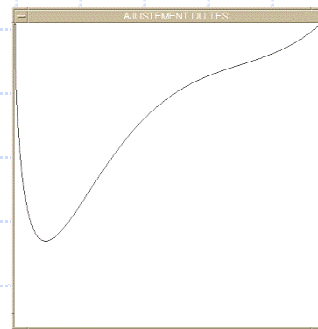
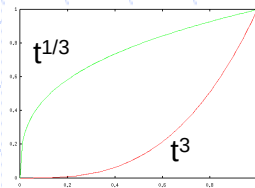
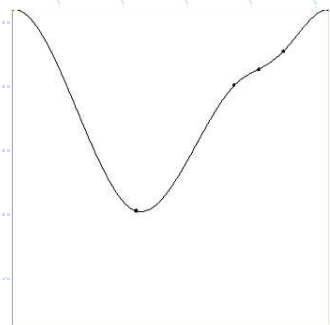


Validation

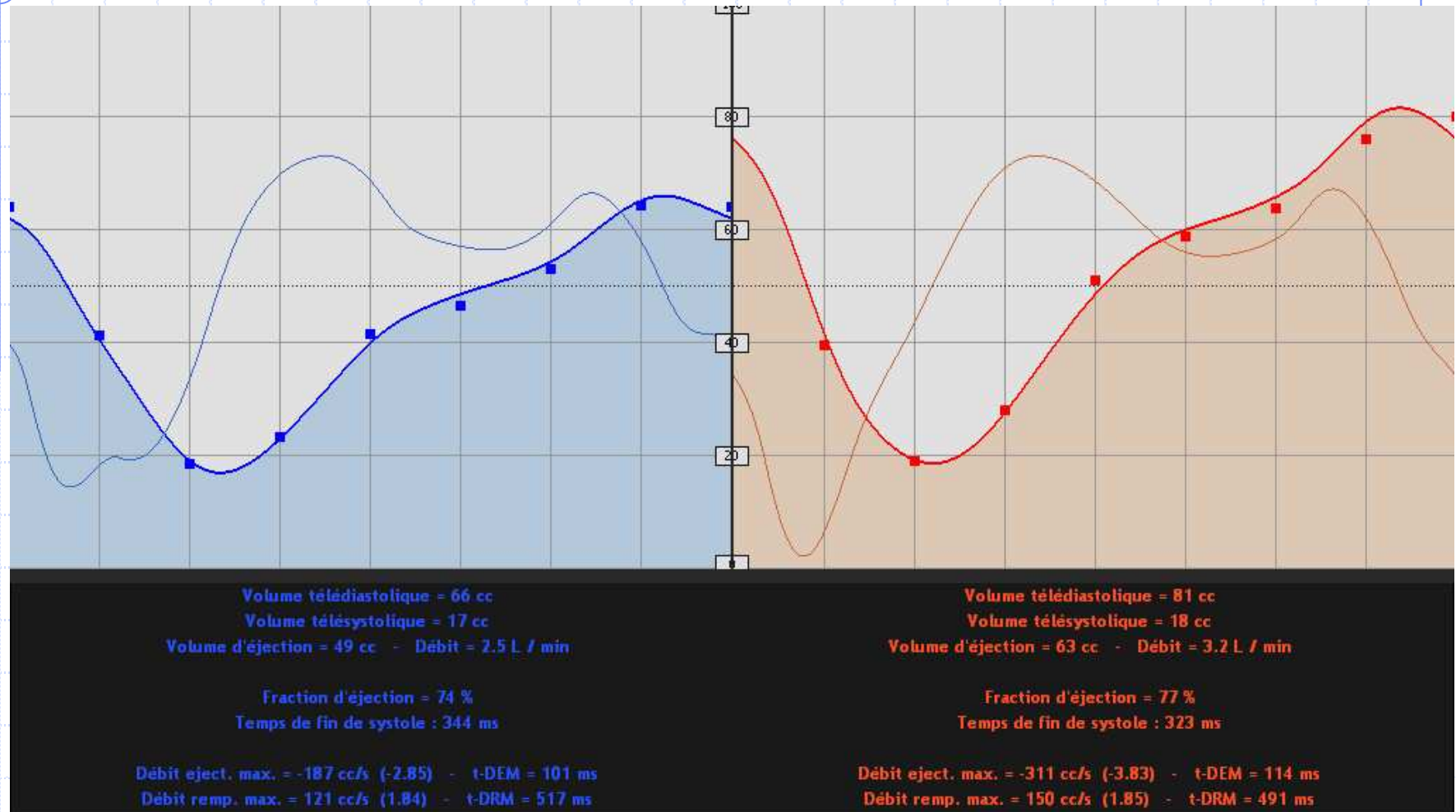
◆ Filtrage linéaire de la CTA

- H1 : imprécis
- Multi-harmoniques : bruité

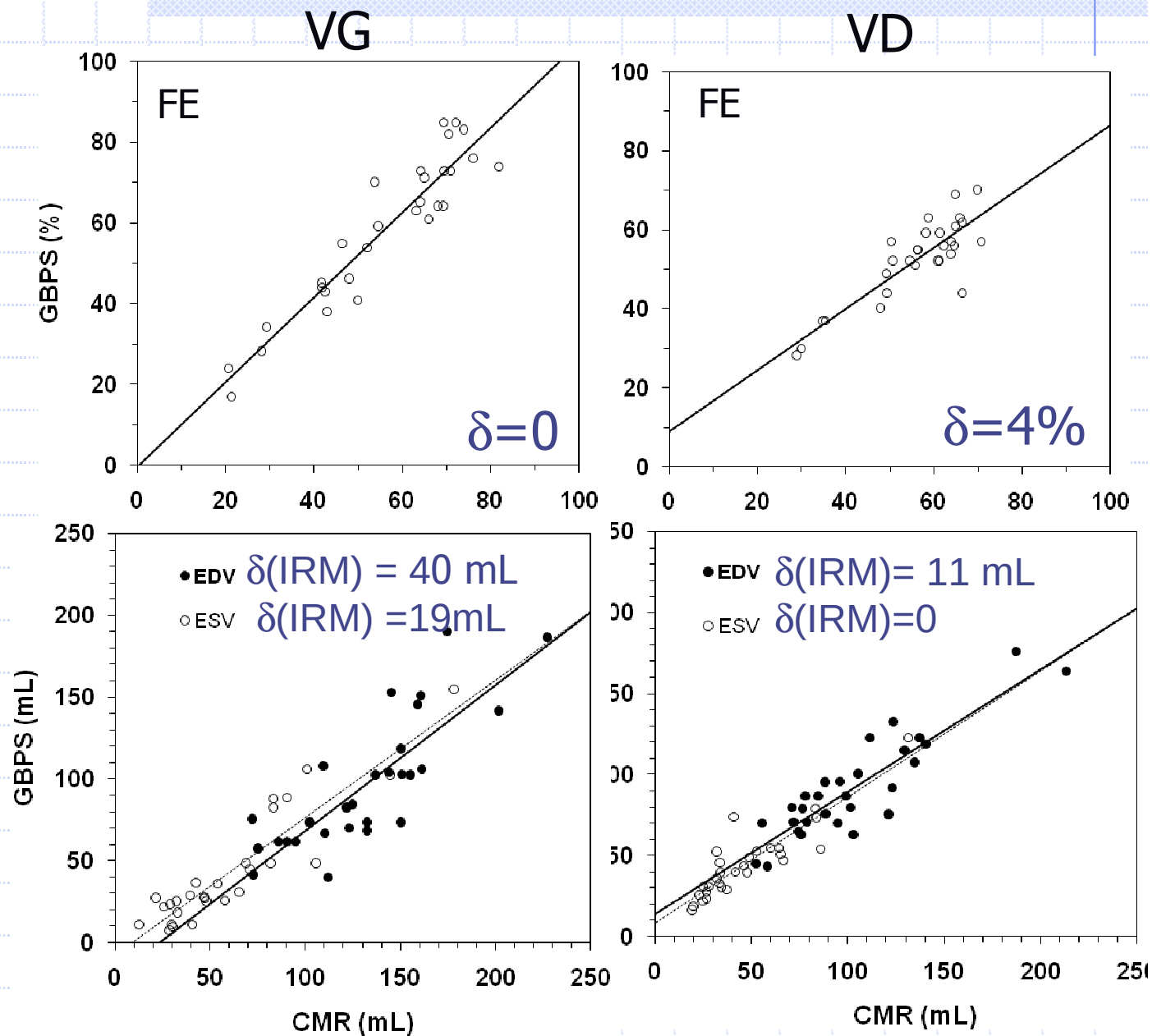
◆ Modélisation de la CTA



Analyse de la fonction systolique

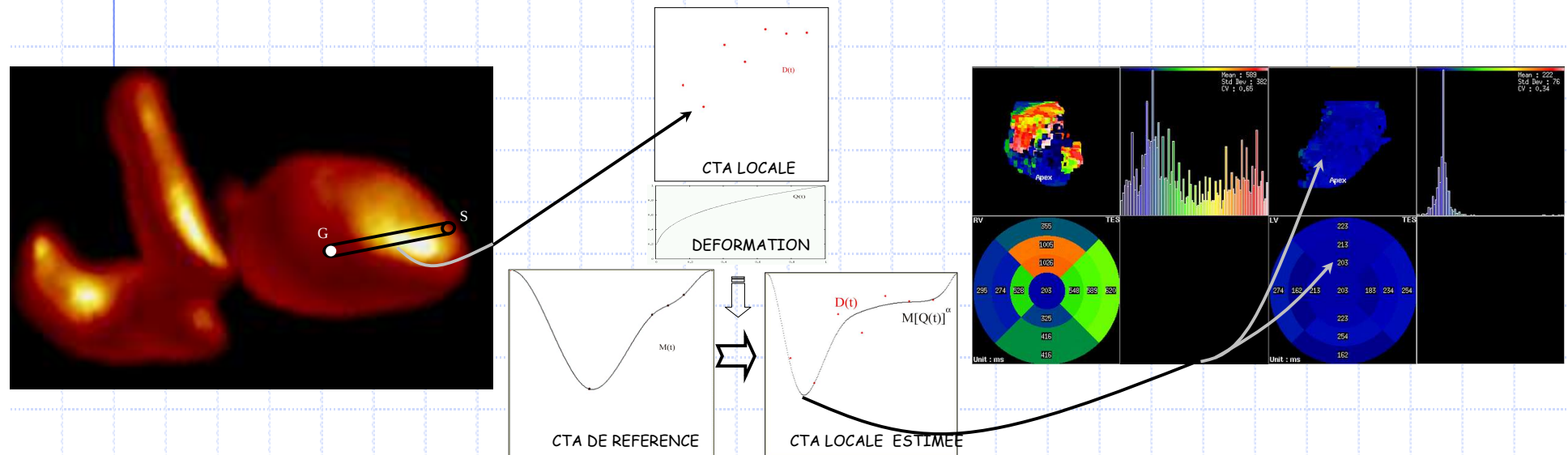


TVI versus IRM

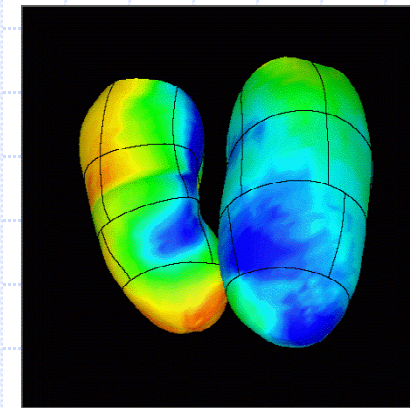
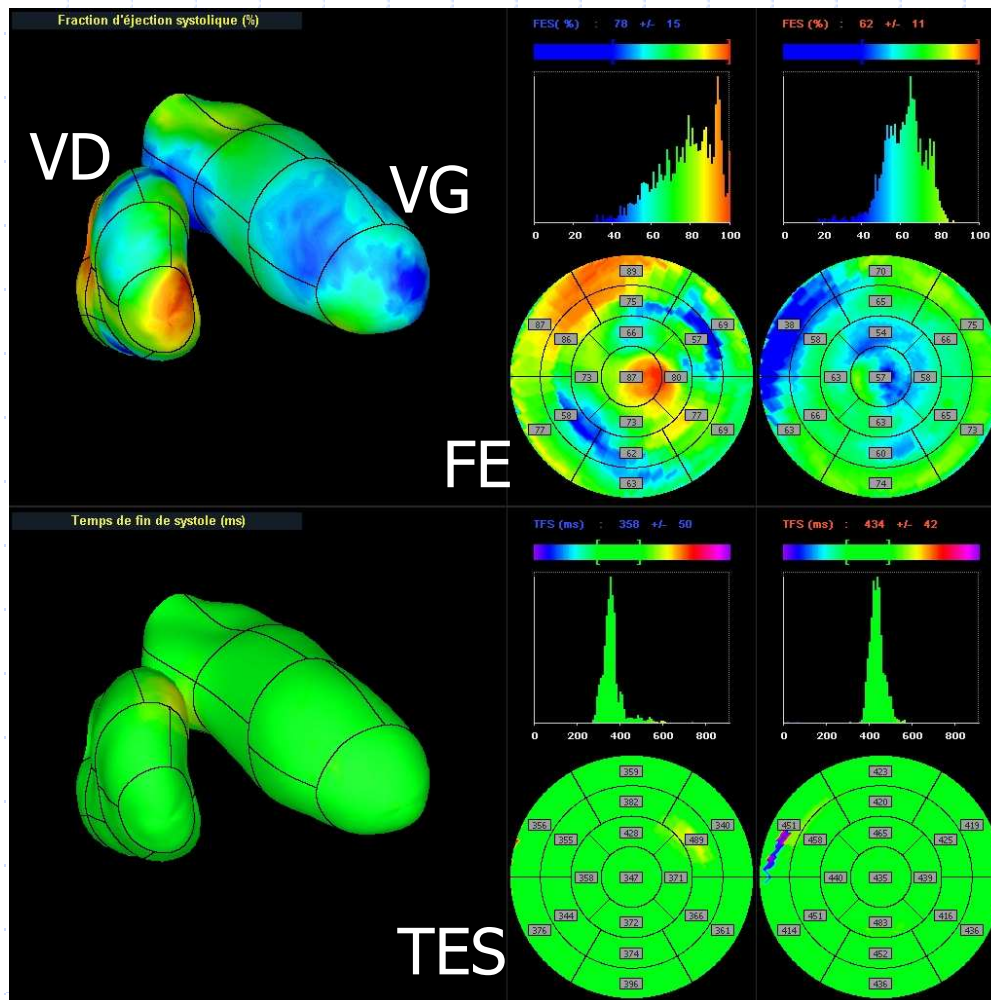


VES(G-D): 9 ± 14 (GBPS) versus 18 ± 13 (IRM)

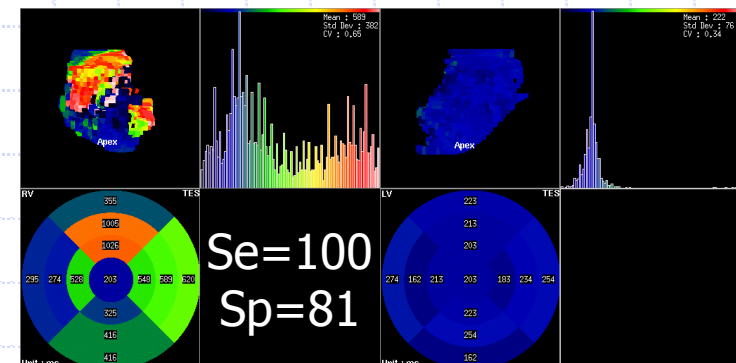
ANALYSE 3D DE CTA LOCALES



ANALYSE 3D DE CTA LOCALES

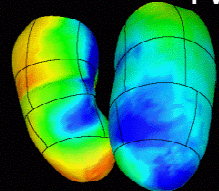
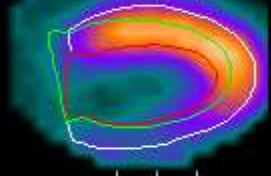
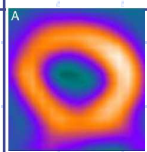
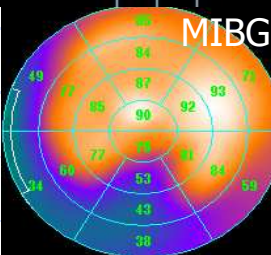
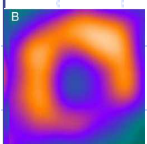


Parameter	Control subjects	Patients with localized ARVD	Patients with diffuse ARVD
EF (%)	63 ± 7	60 ± 5	33 ± 12 [†]
EDV (mL)	101 ± 17	107 ± 20	180 ± 81 [†]
ESV (mL)	37 ± 8	43 ± 10	124 ± 70 [†]
σ-EF (%)	13 ± 3	14 ± 2	11 ± 3
σ-TES (ms)	63 ± 24	167 ± 64 [†]	277 ± 106 [†]



S=80 ms

Synthèse et résultats

Modalité	Logiciels Déconvolution Reconstruction +	FE	Volumes	Cicatrice	Hypo-kinésie	Dys-kinésie
 TVI	LPE Fit de CTA Échantillonnage	+	+		+	+
 32 PERFUSION	Identification Modélisation Quantification	Pronostic		 A	+	+
 MIBG	Identification Modélisation Quantification	-	-	 B	+	-

Pour aller plus loin...

✓ The Mathematics of Computerized Tomography.

F. Natterer. 2001. SIAM.

✓ Quantitative Analysis in Nuclear Medicine Imaging.

H. Zaidi Ed. Springer.

(<http://www.springerlink.com/content/k513k6/?p=be16e74068244dbfbbeb4d7123ee136b3&pi=0>)

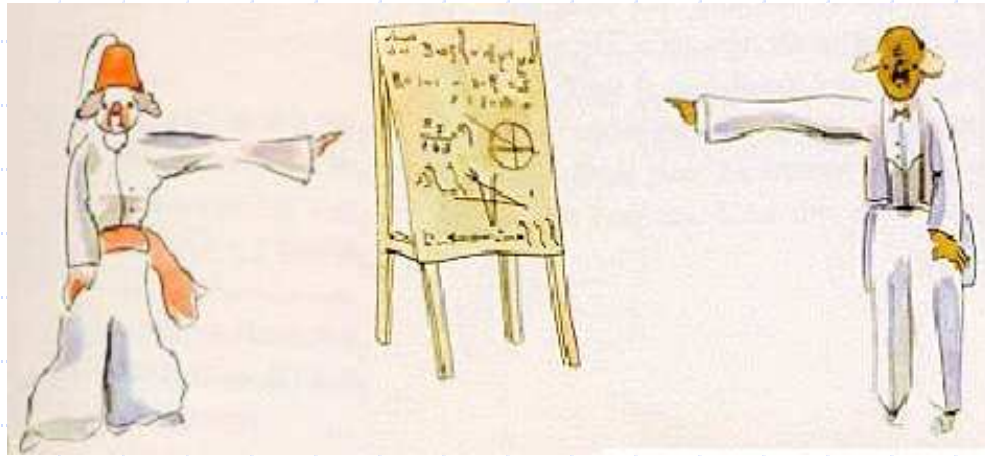
✓ Introduction au traitement numérique des images.

D. Mariano-Goulart. Encyclopédie Médico-chirurgicale, 35-100-A-10, 2009.

✓ Reconstruction tomographique en imagerie médicale.

D. Mariano-Goulart. Encyclopédie Médico-chirurgicale, 35-105-A-10, 2009.

✓ <http://scinti.etud.univ-montp1.fr>



Merci pour votre attention...

d-mariano_goulart@chu-montpellier.fr
<http://scinti.etud.univ-montp1.fr>