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Imagerie Nucléaire, méthode de référence?

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Your Choices in Heart Scans

Once the stethoscope was the only tool doctors had for diagnosing heart trouble. Today, using the latest X-ray, magnetic and ultrasound scanners, they can take detailed, three-dimensional and even moving pictures of that vital organ. Here's a guide to the many choices available. —By Alan Park and Gordon Siskel

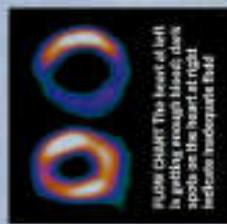
ELECTROCARDIOGRAM (ECG)

- How it works This oldest and most basic heart scan records the electrical impulses that regulate your heart's pumping action. It may seem straightforward, but any disruption from the normal rhythm pattern can alert doctors to the likelihood of enlarged heart tissue and reduced blood flow.
- Limitations While it can pick up signs of trouble, an ECG measures no visual view of the heart and cannot identify exactly what ails the organ or where in the heart the problem lies.



NUCLEAR STRESS TESTING

- How it works Doctors inject a radioactive substance into the blood, then use gamma-ray cameras to see how the blood moves through the heart. The test shows how well the heart is doing at keeping itself saturated with oxygen-rich blood. The test is often done twice, to check cardiac performance at rest and under physical stress.
- Limitations Carrying out two scans can take an hour or two hours. The test also reduces the patient to small amounts of radiation.



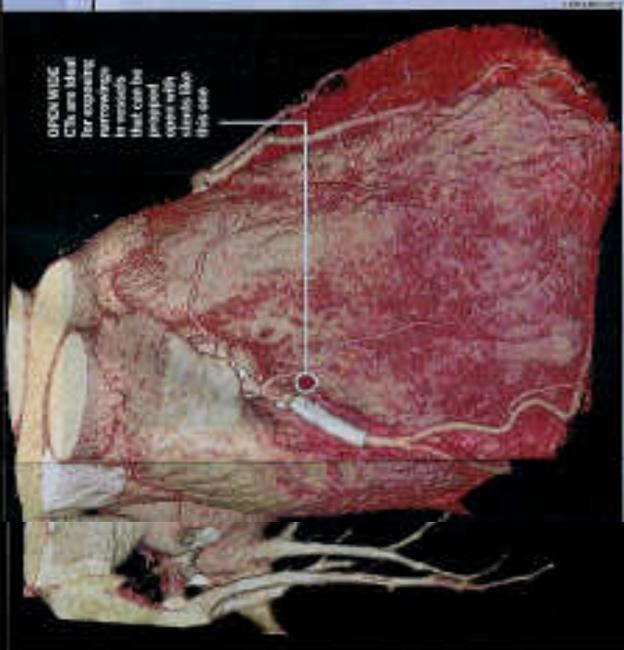
LOW FLOW? The heart at left is getting enough blood; dark spots on the heart at right indicate inadequate flow.

ECHOCARDIOGRAM (ECHO)

- How it works Ultrasonic ultrasound waves, similar to the ones used to take sonograms of a fetus, are directed at the chest and bounce off the heart's walls and valves. A computer analyzes these returning echoes and calculates the size, shape and movement of structures inside the heart. Doctors often take two echoes—one of the heart at rest and another at the heart's widest.



LEAKY SEAL An echo reflects a valve (arrow) that doesn't shut properly, which can reduce the chamber of the heart.



OPEN VEINS CAs are ideal for exposing narrowed or blocked arteries that can be bypassed with open vein grafts like this one.

CT SCAN

- How it works This fast, contrasted rapid X-ray scanner with multiple computerized tomography (CT) to produce the most detailed images available of the heart's arteries without surgery. Patients receive an injection of contrast dye to highlight the blood vessels, and X-rays create images of the heart in slices. A computer generates the slices as an image of the heart that reveals clogged and/or filled plaques lodged in the arteries.
- Limitations CT scans involve radiation exposure, a particular concern for children. Those who are overweight or have asthma or asthma-related calcium deposits won't generate useful images, even for contrast. X-rays and the lasers cannot penetrate metal or calcium.



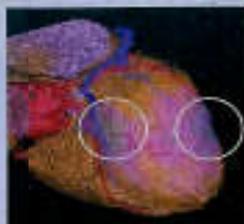
CALCIUM PLUS Bypassing bypasses along the arteries (circled) indicates associated calcium plaques that appear as white on CT scans.



STENTS A powerful CT scanner provides a detailed look at two vein grafts (arrows) inserted to divert blood around arterial blockages.

PET/CT SCAN

- How it works A blend of positron emission tomography and CT, this scan provides structural and functional information about the heart in a single scan. Doctors use the CT to identify blockages, measure plaque along arteries, then apply PET to address parts of the heart muscle, such as the artery linings, that are deprived of blood flow as a result.
- Limitations PET technology is expensive, and the lipid linings are relatively invisible. The scan also involves some radiation exposure.



MAGNETIC RESONANCE IMAGING (MRI)

- How it works Powerful magnets create a field that sets the nuclei of atoms in heart cells vibrating. The resulting atoms emit radio signals, which are converted by a computer into either still or moving 3-D images. The atom at left points to a plaque that's set in the artery; the scan also reveals the layer at top that involves most hearts.
- Limitations Because of the intense magnetic field, patients with pacemakers, stents or other metal implants can't get an MRI. Those scans cannot pick up calcium deposits, which could signal dangerously narrowed arteries.



MRIs can show plaque in arteries.

CORONARY ANGIOGRAM

- How it works This procedure is the gold standard for seeing the arteries that nourish the heart. Doctors insert a catheter through an artery in the leg and shove it up toward the heart. They then send a special dye through the line that highlights the arteries under X-rays and exposes any blockages.
- Limitations Because they're catheters, angiograms have some risks: patients can lose artery walls, require surgical repair. In 2% of cases, serious complications, including death, may occur. Moreover, patients need to be still for four to six hours until the dye is out of the leg vein.



BLOCKED BUREAU A catheter (arrow) that would inject blood flow into the rest of the heart.

- How it works After the patient awakes on a bench or other machines have impacted a dye to make the heart react. Compression of the two images helps pinpoint abnormal valves or areas that are not receiving enough blood.
- Limitations Ultrasound does not produce images with high enough resolution to see heart arteries and can highlight only the biggest changes in structures like the heart's chambers.

Les challengers à la médecine nucléaire

- techniques

- Echocardiographie
- TDM-MB
- IRM

- objectifs

- Perfusion myocardique
- Fonction VG
- Viabilité myocardique

Détection ischémie myocardique en aval sténose coronaire. TSMP effort

Table 1. Diagnostic accuracy of exercise MPS for the detection of $\geq 50\%$ coronary stenosis defined angiographically

Author, year [ref.]	No.	Tracer	Analysis	MI excluded	Quality	Sens (%)	Spec (%)	Acc (%)
Tamaki, 1984 [11]	104	^{201}Tl	Q	Both ^a	2	98	91	96
DePasquale, 1988 [12]	210	^{201}Tl	Q	Both ^a	2	95	74	92
Fintel, 1989 [13]	135	^{201}Tl	V	No	2	83	–	–
Iskandrian, 1989 [14]	193	^{201}Tl	V	Yes	2	86	62	79
Maddahi, 1989 [15]	110	^{201}Tl	Q	Both ^a	3	95	56	88
Mahmarian, 1990 [16]	296	^{201}Tl	Q	Both ^a	3	87	87	87
Van Train, 1990 [17]	242	^{201}Tl	Q	Both ^a	2	94	43	85
Coyne, 1991 [18]	100	^{201}Tl	V	No	2	81	74	77
Quinones, 1992 [19]	112	^{201}Tl	V	Yes	1	77	81	78
Chae, 1993 [20]	243	^{201}Tl	V	No	3	71	63	69
Grover-McKay, 1994 [21]	18	^{201}Tl	V	No	1	91	86	89
Tamaki, 1994 [22]	25	^{201}Tl	V	No	1	95	33	88
Ho, 1997 [23]	51	^{201}Tl	V	Both ^a	3	76	77	76
Kiat, 1990 [24]	53	MIBI	Q	Both ^a	2	94	80	92
Pozzoli, 1991 [25]	75	MIBI	V	No	1	84	88	85
Solot, 1993 [26]	78	MIBI	V	No	2	96	74	90
Marwick, 1994 [27]	86	MIBI	V	Yes	2	73	70	72
Van Train, 1994 [28]	124	MIBI	Q	Both ^a	2	89	36	81
Tamaki, 1994 [22]	26	Tetro	V	No	1	96	67	92
Heo, 1994 [29]	23	Tetro	V	No	2	87	–	–
Benoit, 1996 [30]	30	Tetro	V	Yes	2	81	89	83
Shanoudy, 1998 [31]	26	Tetro	V	No	2	96	–	–

Détection ischémie myocardique en aval sténose coronaire. TSMP dipyridamole

Table 2. Diagnostic accuracy of dipyridamole MPS for the detection of $\geq 50\%$ (or $\geq 70\%^a$) coronary stenosis defined angiographically

Author, year [ref.]	No.	Tracer	Analysis	MI excluded	Quality	Sens (%)	Spec (%)	Acc (%)
Francisco, 1982 [32] ^a	75	²⁰¹ Tl	Q	No	2	90	96	92
Huikuri, 1988 [33] ^a	93	²⁰¹ Tl	V	No	2	96	75	94
Go, 1990 [34]	202	²⁰¹ Tl	V	Both ^b	3	76	80	77
Mendelson, 1992 [35] ^a	79	²⁰¹ Tl	V	Both ^b	2	90	–	–
Cramer, 1994 [36]	38	²⁰¹ Tl	V	No	3	90	71	87
Grover-McKay, 1994 [21]	18	²⁰¹ Tl	V	Unknown	1	91	100	94
Ho, 1995 [37]	54	²⁰¹ Tl	V	No	3	98	73	93
Watanabe, 1997 [38]	53	²⁰¹ Tl	V	Yes	2	80	72	77
Tartagni, 1991 [39]	30	²⁰¹ Tl; MIBI	V	No	2	100	75	97
Miller, 1997 [40]	244	MIBI	V	Both ^b	2	91	28	81
Schillaci, 1997 [41]	40	MIBI	V	Yes	1	95	72	85
Soman, 1997 [42]	27	MIBI	V	No	3	90	66	89
Ogilby, 1998 [43]	26	MIBI	V	No	2	90	100	92
Santoro, 1998 [44] ^a	60	MIBI	V	Yes	3	97	89	93
He, 1997 [45]	64	Tetro	V	No	2	85	54	80

Détection ischémie myocardique en aval sténose coronaire. TSMP adénosine

Table 3. Diagnostic accuracy of adenosine MPS for the detection of $\geq 50\%$ coronary stenosis defined angiographically

Author, year [ref.]	No.	Tracer	Analysis	MI excluded	Quality	Sens (%)	Spec (%)	Acc (%)
Nguyen, 1990 [46]	60	^{201}Tl	V	No	2	92	100	93
Verani, 1990 [47]	45	^{201}Tl	Q	No	2	83	94	87
Coyne, 1991 [18]	100	^{201}Tl	V	Both ^b	2	83	76	79
Nishimura, 1991 [48]	101	^{201}Tl	V	Both ^b	3	84	84	84
Allman, 1992 [49]	76	^{201}Tl	Q	Yes	3	85	38	80
Pennell, 1995 [50]	226	^{201}Tl	V	No	3	96	78	92
Mohiuddin, 1996 [51]	202	^{201}Tl	Q	No	3	90	86	89
Amanullah, 1993 [52]	40	MIBI	V	No	2	94	100	95
Marwick, 1993 [53]	97	MIBI	V	Yes	2	86	71	80
Jamil, 1999 [54]	32	MIBI	V	No	2	75	–	–
Kapur, 2002 [55] ^a	2,560	^{201}Tl , MIBI, Tetro	V	Yes	2	91	87	91

Détection ischémie myocardique en aval sténose coronaire. TSMP dobutamine

Table 4. Diagnostic accuracy of dobutamine MPS for the detection of $\geq 50\%$ (or $\geq 70\%$ ^a) coronary stenosis defined angiographically

Author, year [ref.]	No.	Tracer	Analysis	MI excluded	Quality	Sens (%)	Spec (%)	Acc (%)
Pennell, 1991 [56]	50	²⁰¹ Tl	V	No	2	97	80	94
Warner, 1993 [57]	16	²⁰¹ Tl	V	No	2	93	100	94
Hays, 1993 [58]	67	²⁰¹ Tl	Q	No	2	86	90	87
Huang, 1997 [59]	93	²⁰¹ Tl	Q	No	3	90	81	87
Huang, 1998 [60]	110	²⁰¹ Tl	V	Yes	3	82	82	82
Caner, 1997 [61]	29	²⁰¹ Tl; MIBI	V	Uncertain	3	89	70	83
Gunalp, 1993 [62]	27	MIBI	V	Yes	2	94	89	93
Forster, 1993 [63]	21	MIBI	V	Yes	3	83	89	86
Marwick, 1993 [53]	97	MIBI	V	Yes	2	80	74	77
Marwick, 1993 [64]	217	MIBI	Q	Yes	3	76	67	73
Mairesse, 1994 [65]	129	MIBI	V	Yes	3	76	65	72
Marwick, 1994 [27]	82	MIBI	V	Yes	2	65	68	66
Senior, 1994 [66]	61	MIBI	V	No	3	95	71	88
Di Bello, 1996 [67]	45	MIBI	V	Yes	3	87	86	87
Ifthihar, 1996 [68]	38	MIBI	V	No	2	79	90	82
Kisacik, 1996 [69]	69	MIBI	V	No	3	96	64	86
Slavich, 1996 [70]	46	MIBI	V	Yes	?	82	83	83
San Roman, 1998 [71]	92	MIBI	Q	Yes	3	87	70	82
Santoro, 1998 [44] ^a	60	MIBI	Q	Yes	3	91	81	87
Elhendy, 1998 [72]	70	MIBI	V	No	?	64	72	67

Valeur Pronostique de la TSMP coronariens connus (1)

Table 6. Prognostic value of MPS in definite or suspected CHD (adapted from reference [111])

Year	Author [ref.]	No.	Agent	Abnormal MPS (%)	Mean F/U (m)	HE (%/yr)	HE with abnormal MPS (%/yr)	HE with normal MPS (%/yr)	RR
2001	Galassi [113]	459	Tetro	77	37	2.5	3.0	0.9	3.25
1999	Vanzetto [114]	1,137	²⁰¹ Tl	66	72	1.5	2.0	0.6	3.53
1998	Hachamovitch [108]	5,183	MIBI/ ²⁰¹ Tl	43	21.4	3.0	5.9	0.8	7.50
1998	Olmos [115]	225	²⁰¹ Tl	49	44.4	1.8	2.7	0.9	2.86
1998	Alkeylani [116]	1,086	MIBI	62	27.6	3.4	5.0	0.6	8.92
1997	Snader [117]	3,400	²⁰¹ Tl	21	~24	1.6 (ACM)	~3.8 (ACM)	~1.0 (ACM)	3.75
1997	Boyne [118]	229	MIBI	32	19.2	2.2	5.1	0.8	6.23
1997	Geleijnse [119]	392	MIBI	67	22	6.0	8.7	0.8	10.67
1995	Heller [120]	512	MIBI	58	12.8	4.6	6.9	1.3	5.29
1994	Machecourt [121]	1,926	²⁰¹ Tl	63	33	2.0	2.9	0.5	6.23
1994	Kamal [122]	177	²⁰¹ Tl	83	22	4.3	5.2	0	–
1994	Stratmann [123]	534	MIBI	66	13	10.1	14.3	1.6	9.12
1994	Stratmann [124]	521	MIBI	60	13	4.2	6.7	0.5	14.60

ACM, All-cause mortality; HE, hard event (cardiac death or non-fatal MI); RR, relative risk; other abbreviations as defined in Table 1 and list of abbreviations

Valeur Pronostique de la TSMP

Douleurs thoraciques (2)

Table 7. Prognostic value of normal MPS in patients presenting with stable chest pain (adapted from reference [111])

Year	Author [ref.]	No.	Agent	Normal MPS (%)	Mean F/U (months)	HE with normal MPS (% per yr)
2001	Galassi [113]	459	Tetro	23	37	0.9
2000	Groutars [125]	236	Tetro/ ²⁰¹ Tl	100	25	0.4
1999	Gibbons [126]	4,473	²⁰¹ Tl/MIBI	100	36	0.6
1999	Soman [127]	473	MIBI	100	30	0.2
1999	Vanzetto [114]	1,137	²⁰¹ Tl	34	72	0.6
1998	Hachamovitch [108]	5,183	MIBI/ ²⁰¹ Tl	57	21.4	0.8
1998	Olmos [115]	225	²⁰¹ Tl	51	44.4	0.9
1998	Alkeylani [116]	1,086	MIBI	38	27.6	0.6
1997	Snader [117]	3,400	²⁰¹ Tl	79	~24	~1.0 (ACM)
1997	Boyne [118]	229	MIBI	68	19.2	0.8
1997	Geleijnse [119]	392	MIBI	33	22	0.8
1995	Heller [120]	512	MIBI	42	12.8	1.3
1994	Machecourt [121]	1,926	²⁰¹ Tl	37	33	0.5
1994	Kamal [122]	177	²⁰¹ Tl	17	22	0
1994	Stratmann [123]	534	MIBI	34	13	1.6
1994	Stratmann [124]	521	MIBI	40	13	0.5
	Total	20,963		53	28.3	0.7

ACM, All-cause mortality; HE, hard event (cardiac death or non-fatal MI); RR, relative risk; other abbreviations as defined in Table 1 and list of abbreviations

Valeur Pronostique de la TSMP Pré opératoire(3)

Table 2. MPI for preoperative assessment of cardiac risk (adapted from references [15-6])

Year	Author [ref.]	No.	Infarctus ischemia (%)	Mortality (%)	PPV	NPV
Vascular surgery						
1985	Bonchou [133]	46	16 (35)	3 (6%)	19% (3/16)	100% (32/32)
1987	Corder [136]	116	54 (47)	11 (10%)	20% (11/54)	100% (30/30)
1988	Murcho [137]	67	13 (20)	3 (4%)	20% (3/13)	100% (56/56)
1988	Jacobs [138]	46	14 (31)	3 (4%)	14% (3/14)	100% (24/24)
1989	Engle [139]	300	82 (43)	15 (2%)	16% (13/82)	98% (81/82)
1990	Malkous [140]	95	34 (36)	7 (7%)	8% (3/34)	96% (44/46)
1990	Yoniss [141]	111	40 (36)	3 (7%)	13% (6/40)	100% (31/31)
1991	Mangano [142]	60	22 (37)	3 (5%)	9% (1/22)	93% (19/20)
1991	Stevens [143]	68	n/a	4 (6%)	n/a	100% (21/21)
1991	Walton [144]	26	15 (58)	3 (12%)	20% (3/15)	100% (11/11)
1992	Hussel [145]	327	167 (51)	28 (2%)	14% (28/167)	94% (87/98)
1992	Laito [146]	325	161 (49)	30 (2%)	17% (28/161)	94% (180/183)
1992	Martens [147]	68	43 (63)	5 (5%)	11% (5/43)	100% (20/20)
1993	Brown [148]	391	77 (20)	12 (5%)	13% (10/77)	98% (120/121)
1993	Kosowski [149]	170	67 (39)	5 (9%)	4% (3/67)	98% (64/65)
1994	Kono [150]	437	168 (38)	23 (2%)	4% (7/168)	96% (195/203)
1994	Koy [151]	237	116 (49)	17 (7%)	11% (13/116)	100% (87/87)
1995	Kontolau [152]	105	42 (40%)	3 (2%)	6% (3/42)	100% (49/49)
1995	Marshall [153]	117	55 (47%)	12 (10%)	16% (8/55)	97% (32/34)
1997	Van Thromb [154]	142	48 (34%)	3 (2%)	n/a	n/a
Non-vascular surgery						
1990	Camp [155]	40	9 (23)	6 (15%)	67% (6/9)	100% (22/22)
1991	Igbot [156]	31	11 (41)	3 (11%)	27% (3/11)	100% (20/20)
1992	Colby [157]	160	36 (36)	4 (4%)	8% (3/36)	98% (82/84)
1992	Shaw [158]	60	28 (47)	6 (10%)	21% (6/28)	100% (19/19)
1993	Talbot [159]	23	15 (65)	6 (11%)	27% (4/15)	100% (32/32)
1994	Yoniss [160]	161	58 (31)	15 (2%)	18% (8/58)	98% (87/89)
1996	Shintani [161]	229	67 (29%)	10 (4%)	6% (4/67)	94% (81/82)
	Weighted average	3,718		246 (7%)	12.1% (186/1,547)	98.6% (1,744/1,571)

MI, myocardial infarction; NPV, negative predictive value; PPV, positive predictive value; n/a, not available

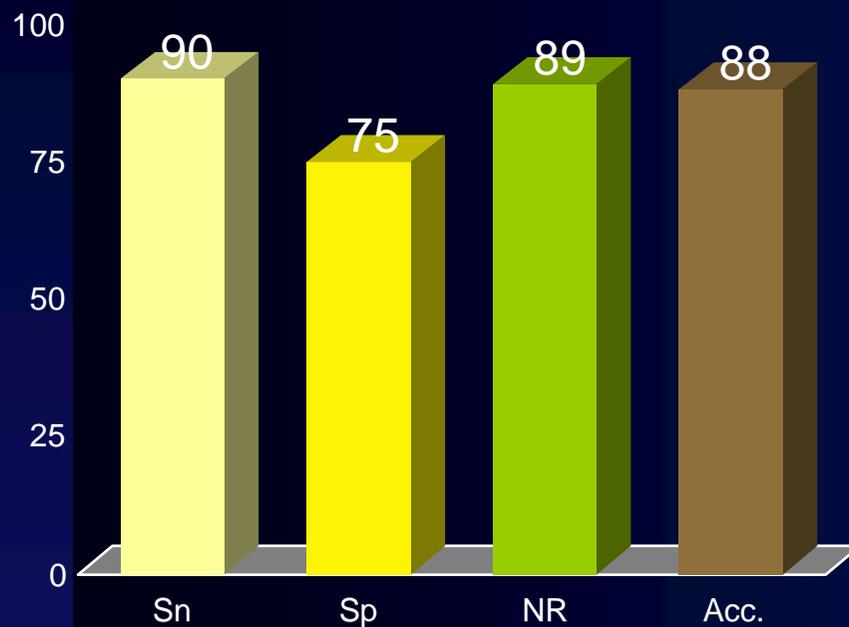
Valeur Pronostique de la TSMP (4)

Diabétique asymptomatique

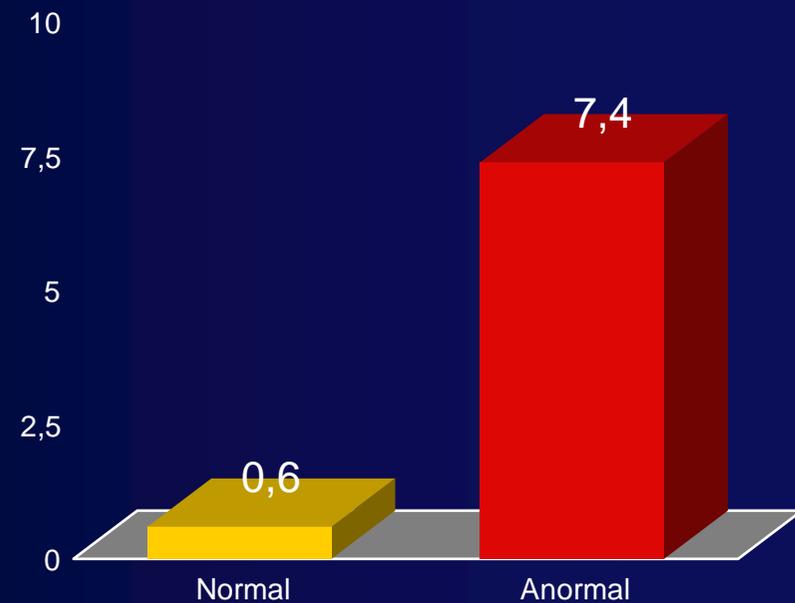
<i>Study</i>	<i>Publications</i>	<i>Yr</i>	<i># pts</i>	<i>F-Up (m)</i>	
Rubler	Am J Cardiol	1987	123	36	
Brown	Am J Cardiol	1989	36	24	Renal failure
Lane	Am J Cardiol	1989	101	< 1	Vascular surg.
Camp	Am J Cardiol	1990	40	< 1	Renal failure
Koistinen	Br Heart J	1990	33	-	
Holley	Am J Med	1991	150	< 1	renal transpl.
Marwick	Transpl	1992	150	-	renal transpl.
Paillole	Diabetologica	1995	59	-	
Vanzetto	Diab Care	1999	158	24	high risk
Kang	Am Heart J	1999	1271	24	
Penfornis	Diabet Med	2001	56	-	versus DES
Kumar	Nucl Med Com	2001	43	-	
Giri	Circulation	2002	929	30	
Faglia	Diabetes care	2002	735	60	
De Lorenzo	Am J Cardiol	2002	180	36	
Berman	JACC	2003	589	27	
MiSAD	Diabetes Care	2003	97	24	
Vanzetto	AHA	2003	756	18	
Miller	Am H J	2004	1738	-	
Pooled data	-	-	7244	36	

Validation Clinique de la TSMP

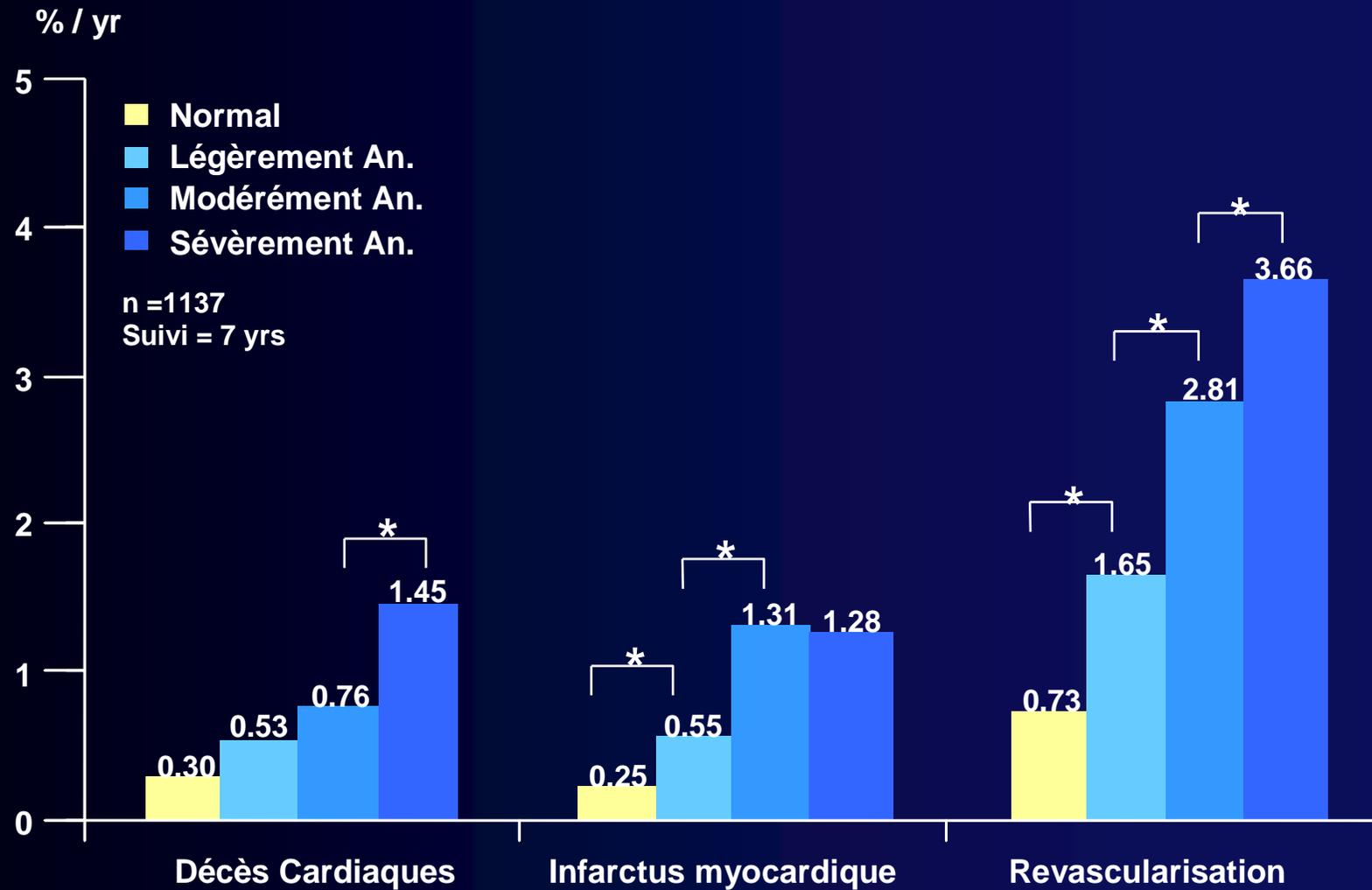
Valeur Diagnostique
79 études - 9000 pts



Valeur Pronostique
60 études - 30 000 pts



Valeur Pronostique de la TSMP (5)



Validation de la TSMP

DTA

PTCA

Sportif

Ponté

Post-IDM

Greffé
cardiaque

Homme

Pré-op

Dysfonction
endothéliale

Femme

Diabétique

Ins.
Rénal

MCH

Ischémie
silencieuse

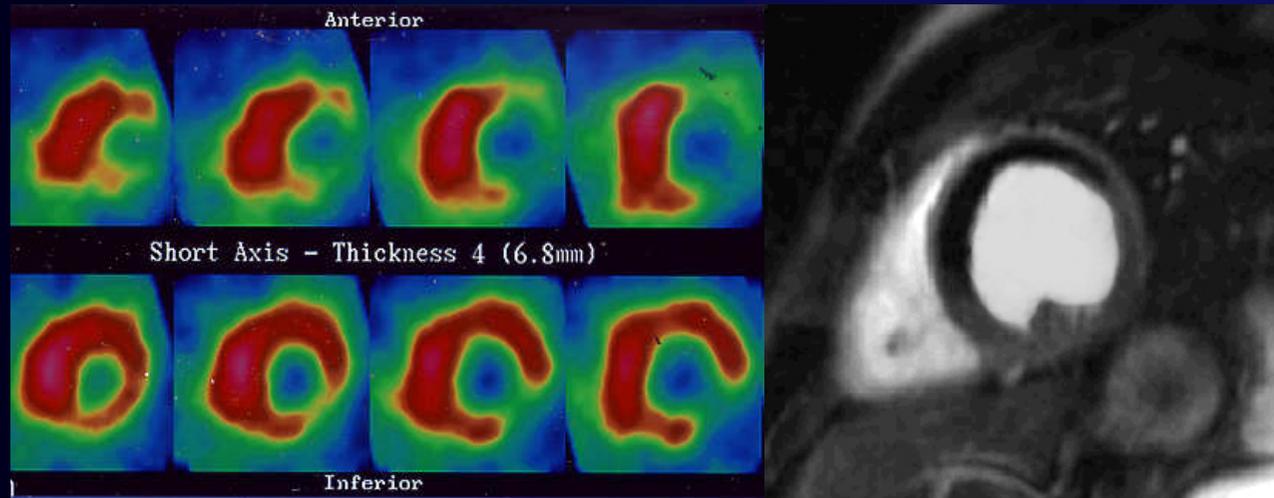
Transplanté
rénal

Vasculaire

ST-SCA

HTA

Ischémie myocardique DES, ECM,IRM, Scanner



Valeur Diagnostique de l'ESD et ischémie

Table 3. Selected Studies Outlining the Accuracy of Exercise Echocardiography*

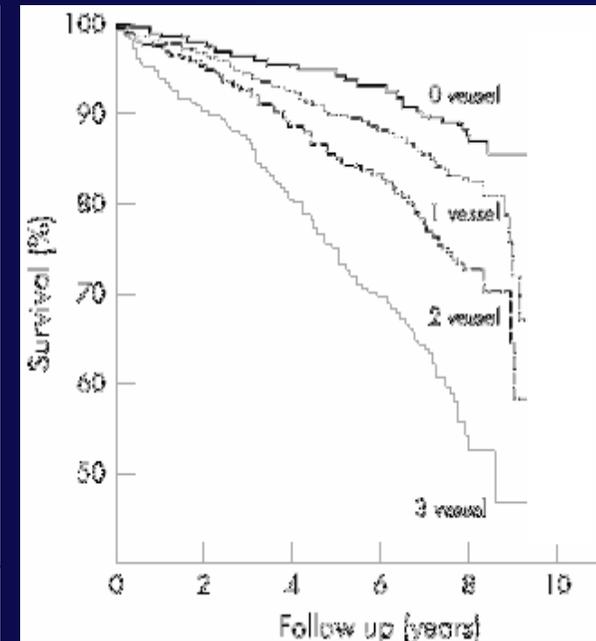
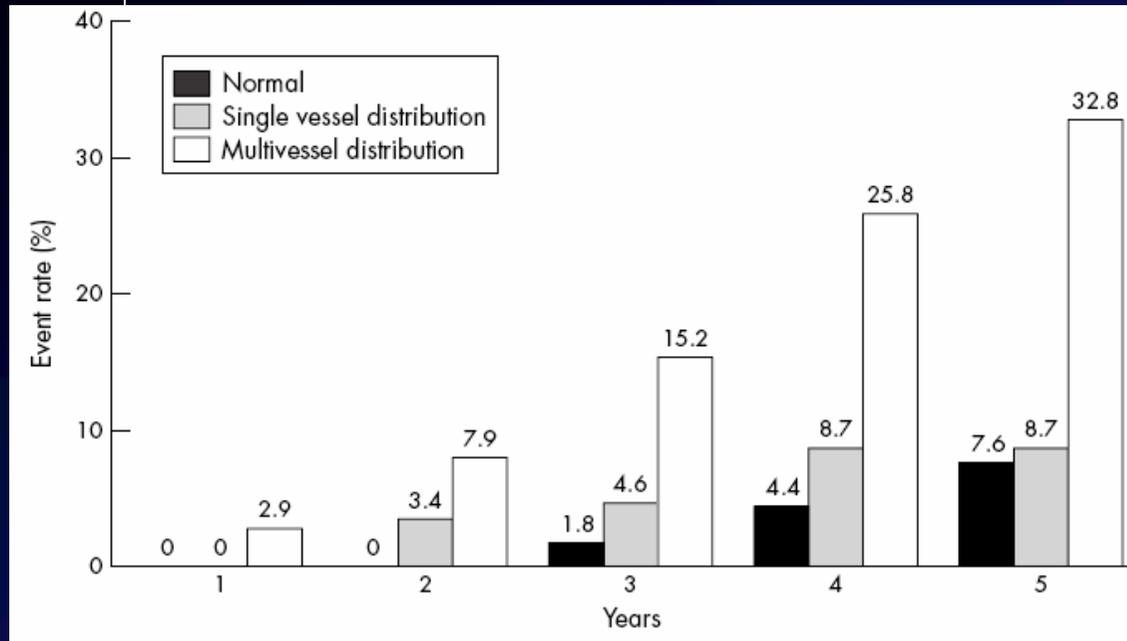
Author (Ref.)	Year	Exercise	Significant CAD	Total No. of Patients	Sensitivity (%)	Sensitivity for 1-VD (%)	Sensitivity for MVD (%)	Specificity (%)	PPV (%)	NPV (%)	Overall Accuracy (%)
Armstrong et al. (4)	1987	TME	50%	123	88	81	93	86	97	61	88
Crouse et al. (5)	1991	TME	50%	228	97	92	100	64	90	87	89
Marwick et al. (6)	1992	TME	50%	150	84	79	96	86	95	63	85
Quinones et al. (7)	1992	TME	50%	112	74	59	89	88	96	51	78
Hecht et al. (8)	1993	SBE	50%	180	93	84	100	86	95	79	91
Roger et al. (9)	1994	TME	50%	150	91	—	—	—	—	—	—
Beleslin et al. (10)	1994	TME	50%	136	88	88	91	82	97	50	88
Roger et al. (11)	1995	TME	50%	127	88	—	—	72	93	60	—
Marwick et al. (12)	1995	TME	50%	161	80	75	85	81	71	91	81
Marwick et al. (13)	1995	TME	>50%	147	71	63	80	91	85	81	82
Luotolahti et al. (14)	1996	UBE	50%	118	94	94	93	70	97	50	92
Roger et al. (15)	1997	TME	50%	340	78	—	—	41	79	40	69
Segar et al. (17)	1992	DSE (5-30)	50%	88	95	—	—	82	94	86	92
Marcovitz et al. (18)	1992	DSE (5-30)	50%	141	96	95	98	66	91	84	89
McNeill et al. (19)	1992	DASE (10-40)	50%	80	70	—	—	88	89	67	78
Marwick et al. (20)	1993	DSE (5-40)	50%	217	72	66	77	83	89	61	76
Previtali et al. (21)	1993	DSE (5-40)	50%	80	79	63	91	83	92	61	80
Takeuchi et al. (22)	1993	DSE (5-30)	50%	120	85	73	97	93	95	80	88

*Modified with permission from the American College of Cardiology Foundation and American Heart Association (2).

DASE = dobutamine-atropine stress echocardiography; DSE = dobutamine stress echocardiography; other abbreviations as in Table 3.

	Exercice	Dobutamine
Etudes	15	28
Patients	1849	2246
Sensibilité	84 %	80 %
Spécificité	82 %	84 %
Accuracy	83 %	82 %

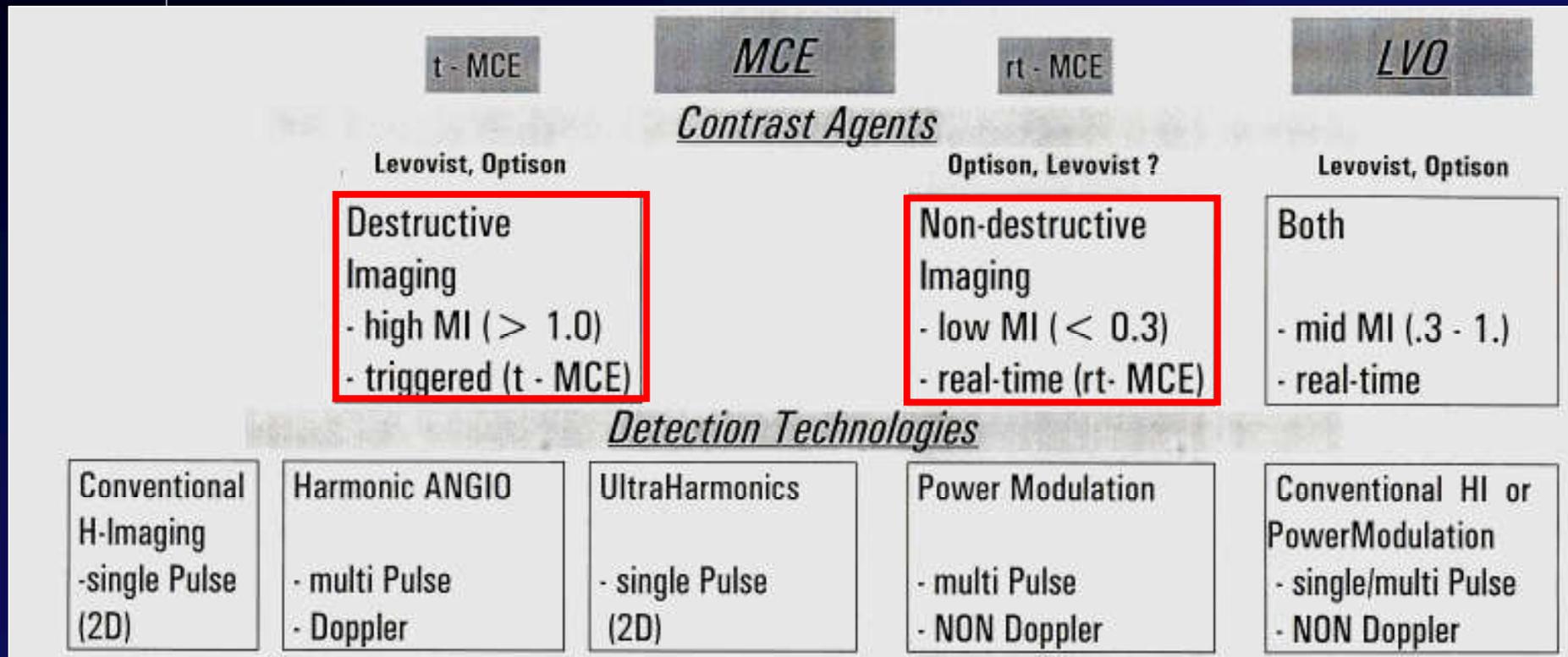
Valeur Pronostique de l'ESD



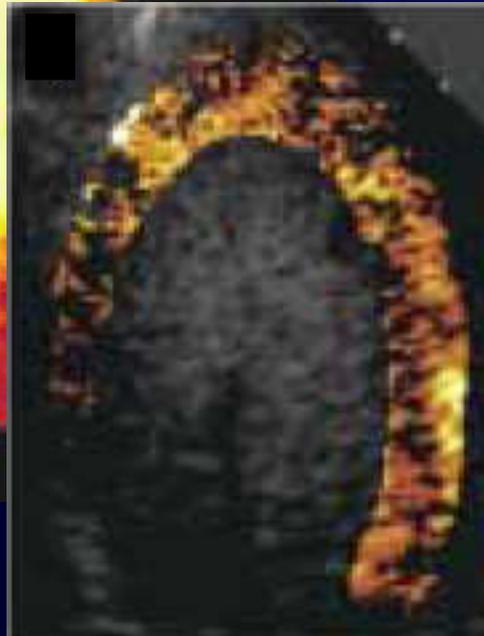
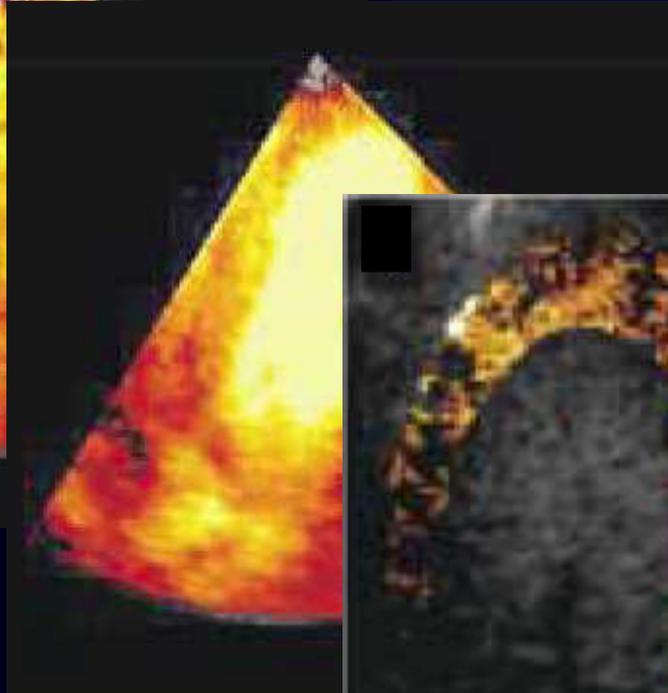
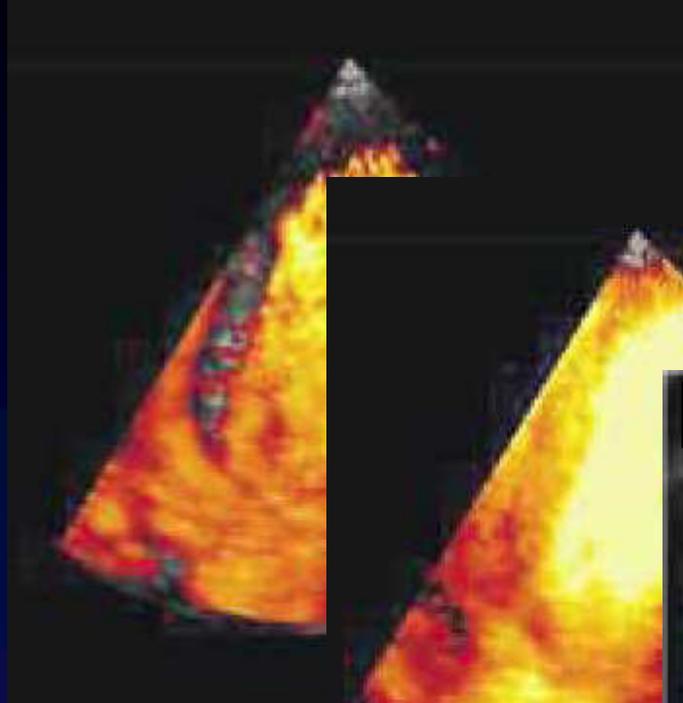
10 Etudes, 27 000 patients
ESD normale : †CV annuelle \pm 1.0% / an
ESD patho : †CV annuelle x 2-5

VPN < isotopes

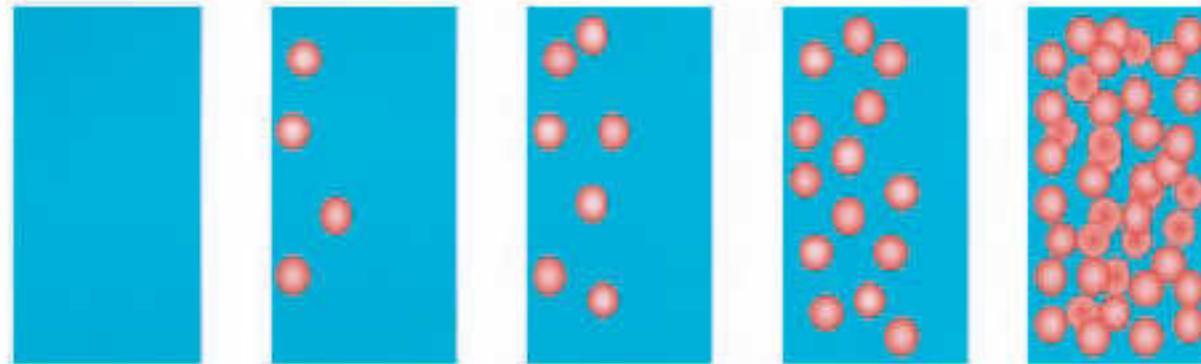
Perfusion : ECM



Perfusion : ECM



Perfusion : ECM



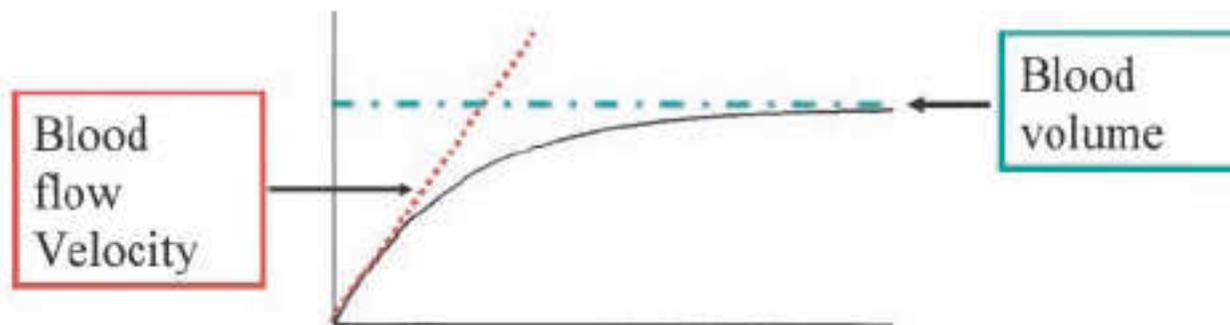
PI=0s

PI=1s

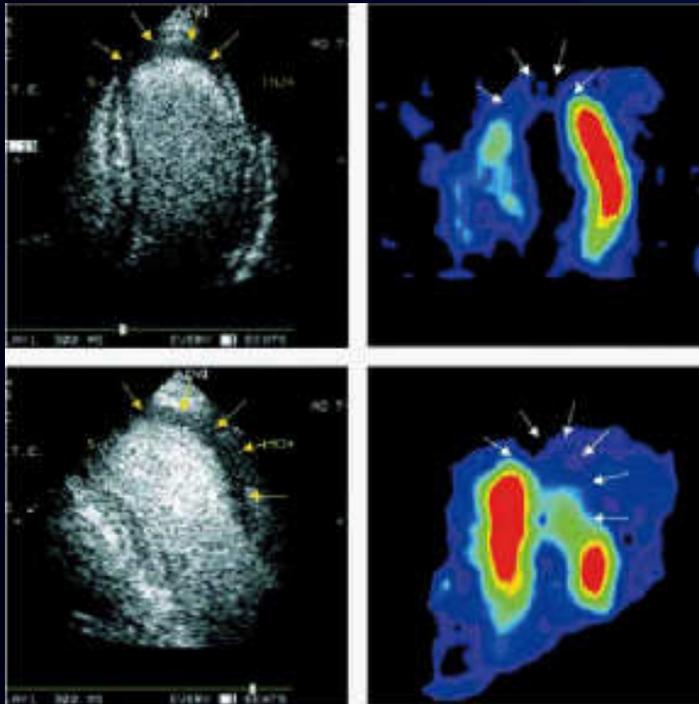
PI=2s

PI=3s

PI=4s



Perfusion : ECM



- 7 études cliniques
- 245 patients
- Comparaison coronarographie
- **Sensibilité 89%**
- **Spécificité 63%**

- **Concordance TSMP**
 - Brute 62%
 - "Corrigée" 82%

Perfusion : ECM

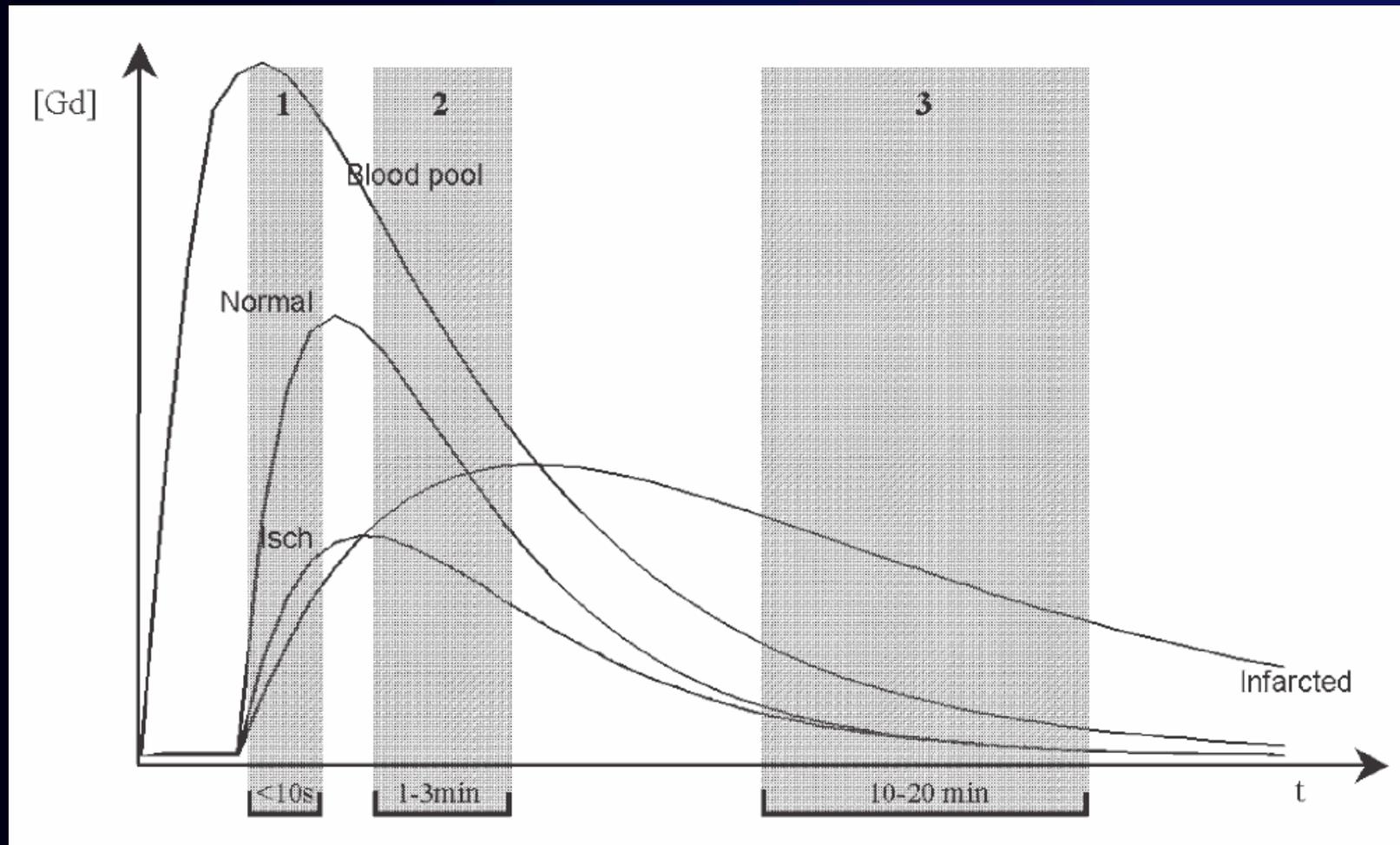
- **Avantages**

- Pas d'irradiation
- Pas d'injection de contraste iodé
- Disponibilité
- Accessible au cardiologue
- Etude anatomique et fonctionnelle cardiaque

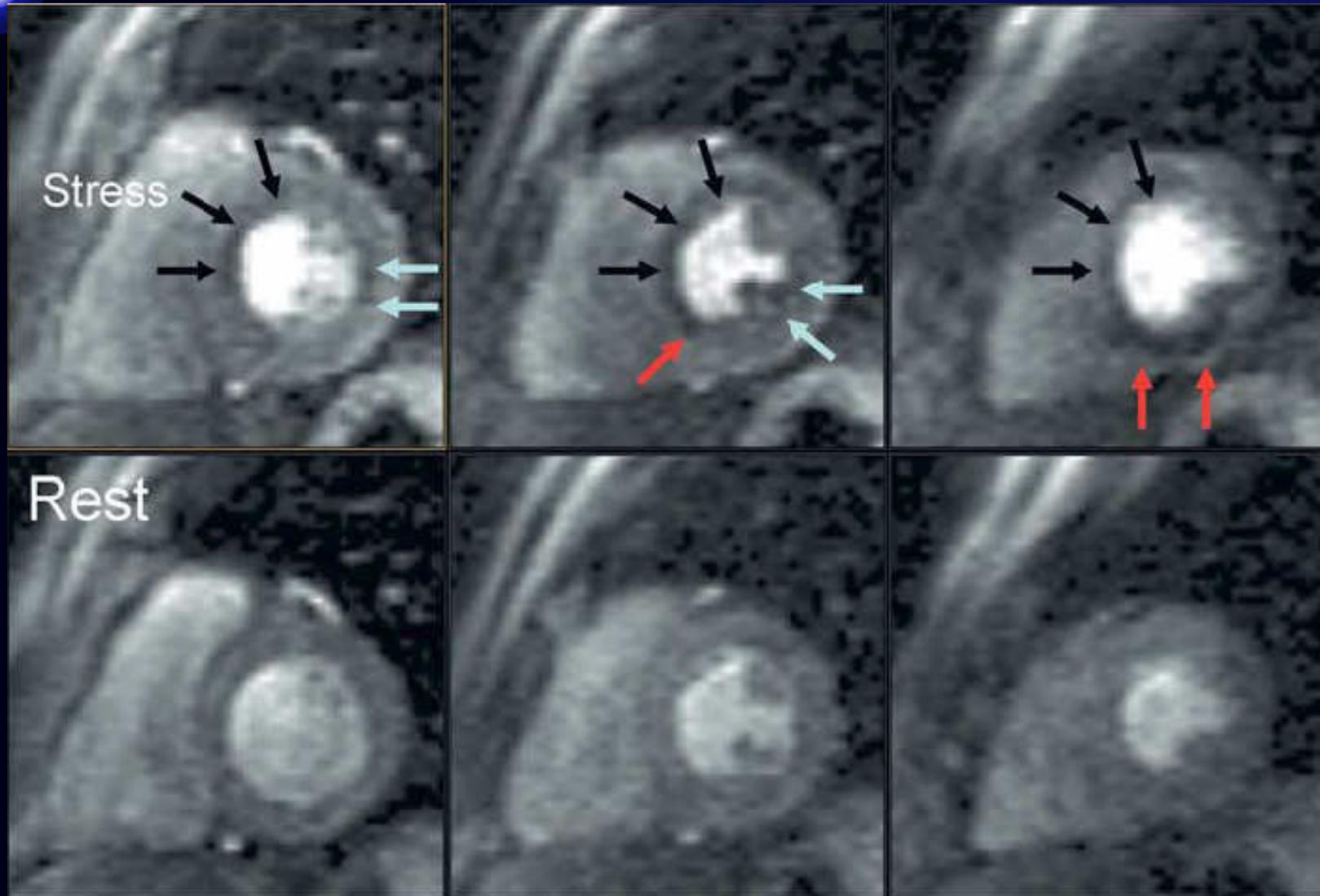
- **Inconvénients**

- Temps médecin (post-traitement)
- Courbe d'apprentissage ++
- Disponibilité des produits de contrastes de 3ème génération
- Echogénicité
- Artéfacts (shadowing, blooming, wall motion artifacts)
- Pas de validation pronostique
- Cotation

Perfusion : IRM Gadolinium



Perfusion : IRM



Perfusion : IRM

- IRM de perfusion et Δc de coronaropathie :
 - 17 études
 - 502 patients (dipyridamole ou adénosine)
 - Sensibilité 84 - 93%
 - Spécificité 63 - 85%

Perfusion : IRM

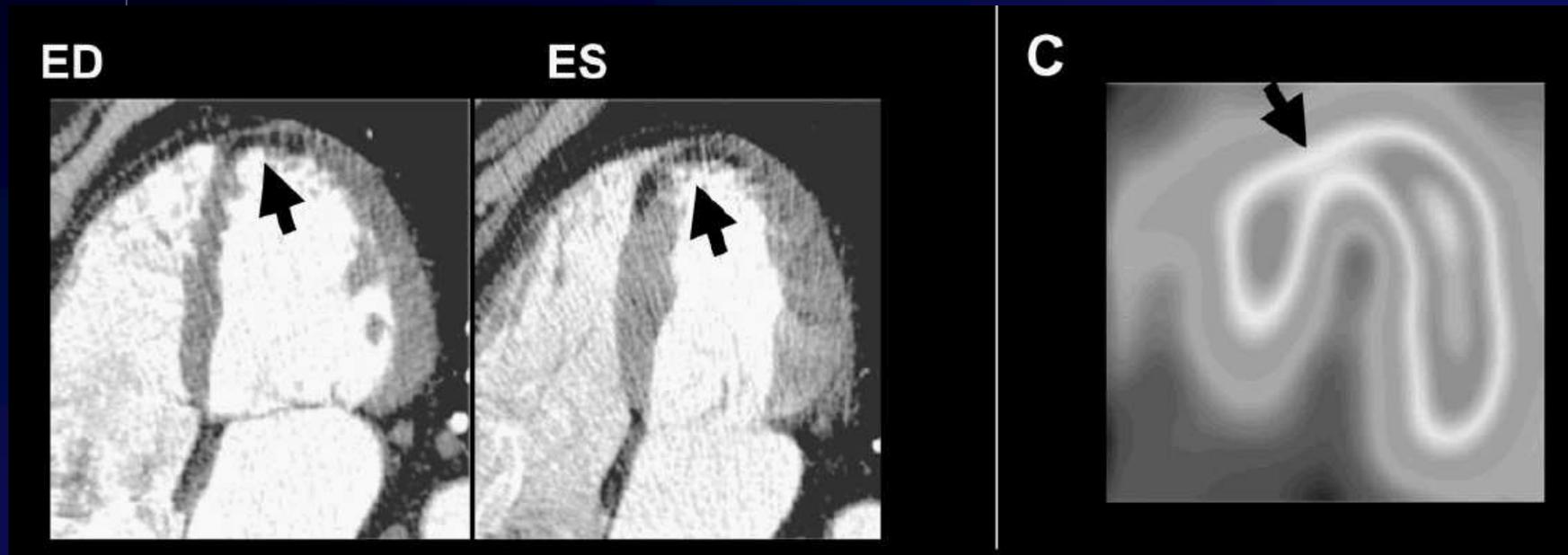
- **Avantages**
 - Pas d'irradiation
 - Pas d'injection de contraste iodé
 - Résolution spatiale
 - Combinaison possible avec IRM dobutamine
- **Inconvénients**
 - Temps médecin (quantification)
 - Disponibilité
 - Courbe d'apprentissage ++
 - Accessibilité au patient
 - Contre-indication aux champs magnétiques
 - Pas de validation pronostique

Perfusion : IRM

"Unfortunately, MRI is still limited to highly specialised centres and acquisition protocols are still time consuming, making the technique currently unsuitable for evaluation of larger populations.

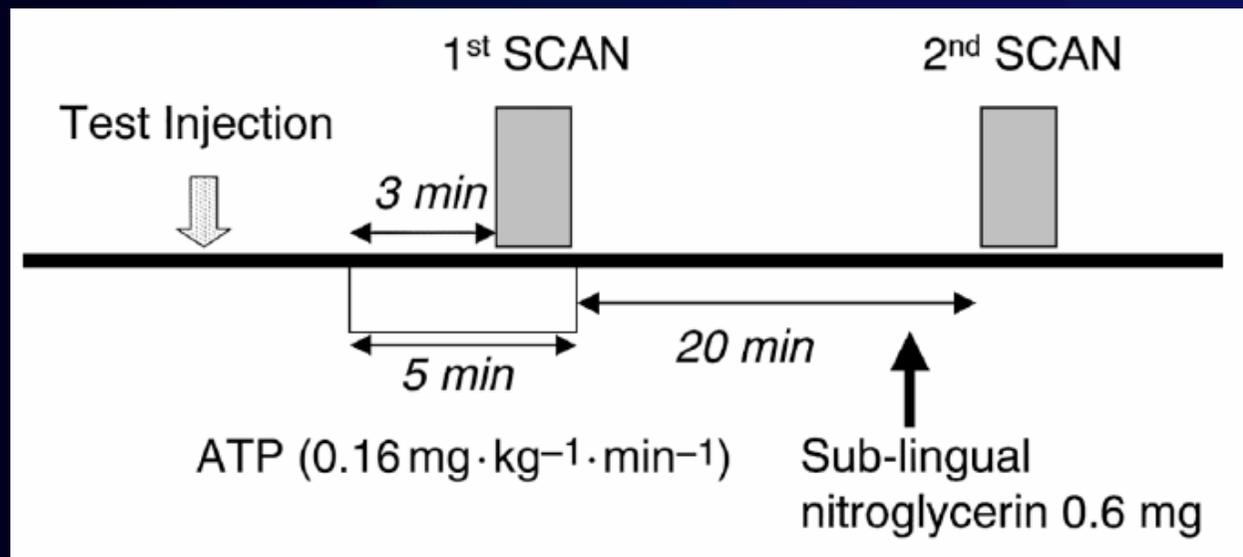
No MRI studies with integration of systolic wall motion and perfusion to detect CAD are currently available."

Perfusion : TDM-MB

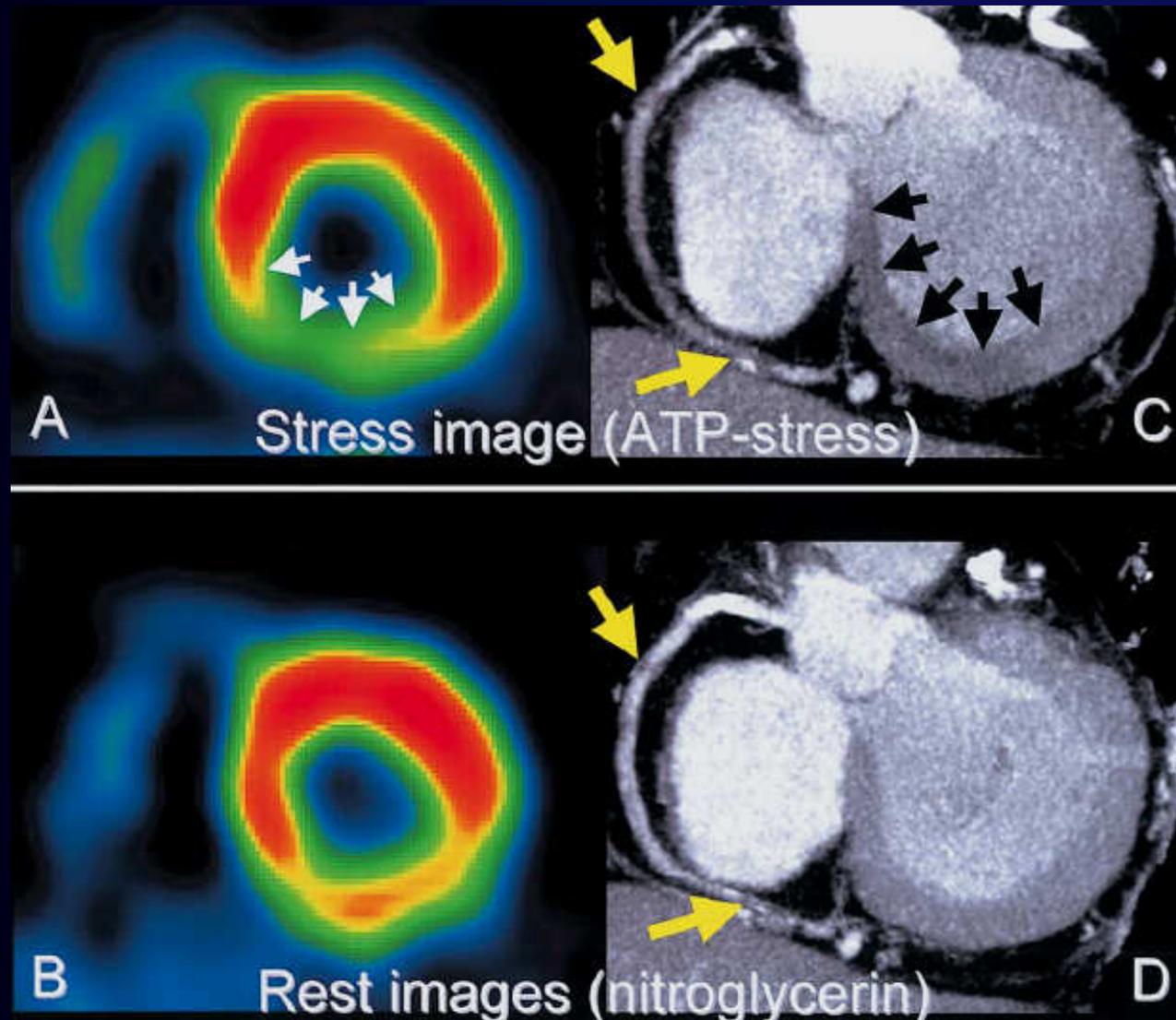


Perfusion : TDM-MB

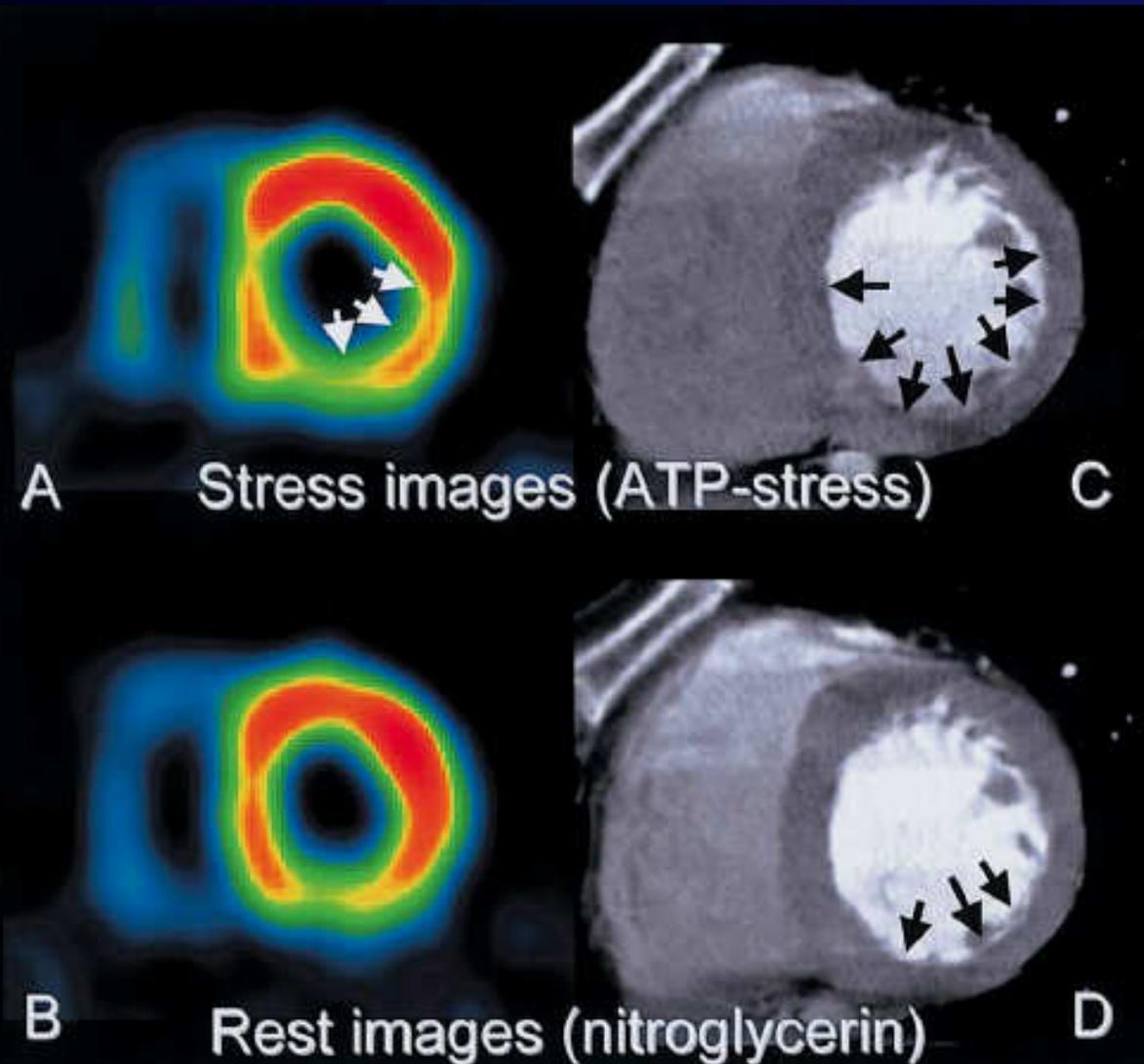
- 84 patients
- TDM 16 barettes
- 50 mg aténolol (< 65 bpm)



Perfusion : TDM-MB



Perfusion : TDM-MB



Perfusion : TDM-MB

- Concordance TDM-MB / TSMP = 83%

Table 1 Agreement Between Perfusion CT and MPS

	<i>MPS positive</i>	<i>MPS negative</i>
<i>Perfusion CT positive</i>	21	5
<i>Perfusion CT negative</i>	1	9

Perfusion CT positive and negative = CE-MSCT described a hypo-perfusion area (HPA) and no-HPA in stress images; MPS positive and negative; stress thallium-201 myocardial perfusion scintigraphy described the presence and absence of redistribution visually noted in the 4-h images. The assessment by perfusion CT was significantly concordant with that by MPS (30/36, $p < 0.05$).

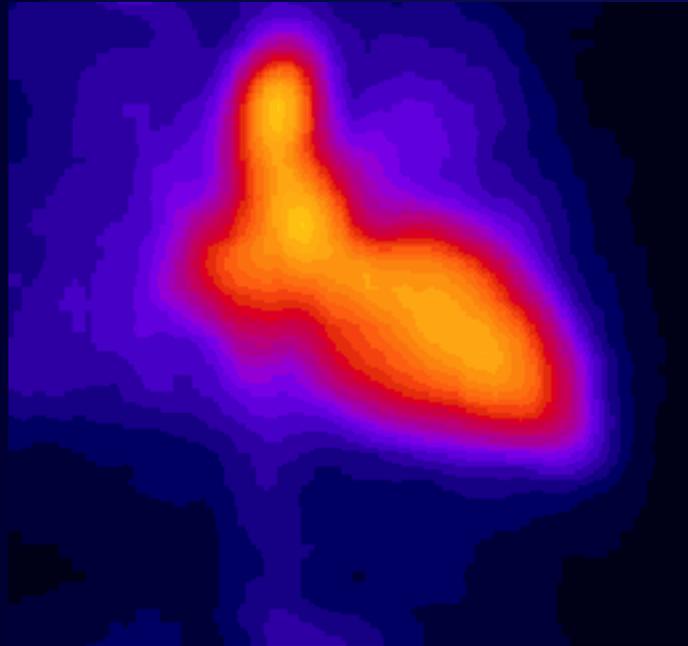
Perfusion : TDM-MB

Our results showed that the ATP stress CE-MSCT can describe both myocardial ischemia induced by the ATP and coronary artery stenosis.

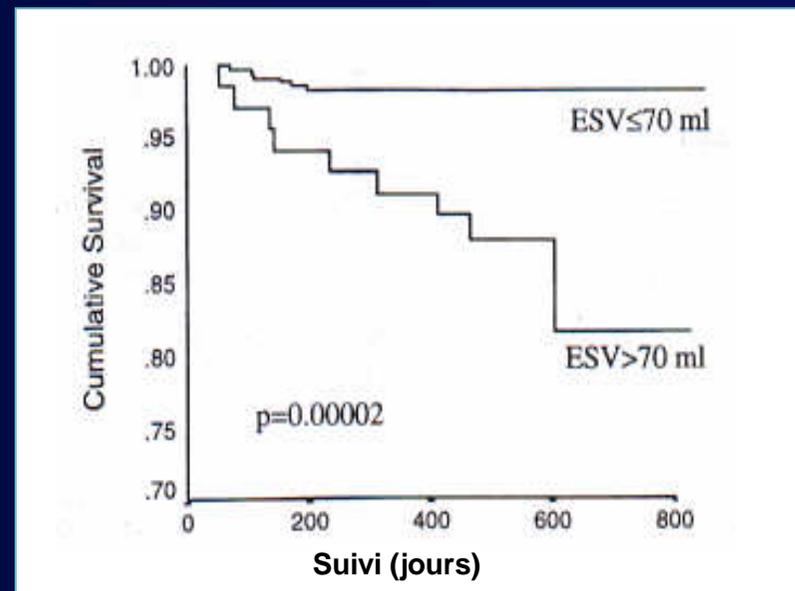
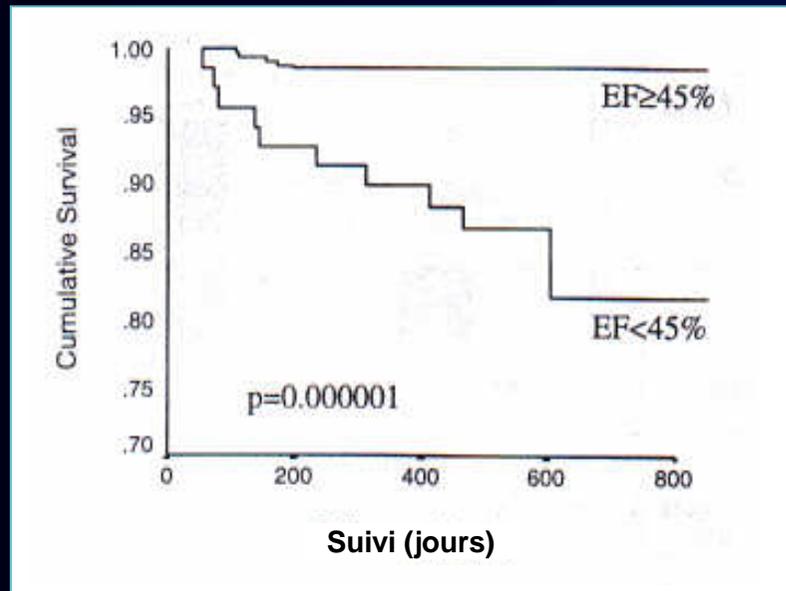
Although the coronary artery stenosis could be assessed with stress images, rest images are of higher image quality and seem to be more feasible at present.

In a clinical setting, ATP stress CE-MSCT is a potential alternative to stress MPS in the evaluation of patients with CAD.

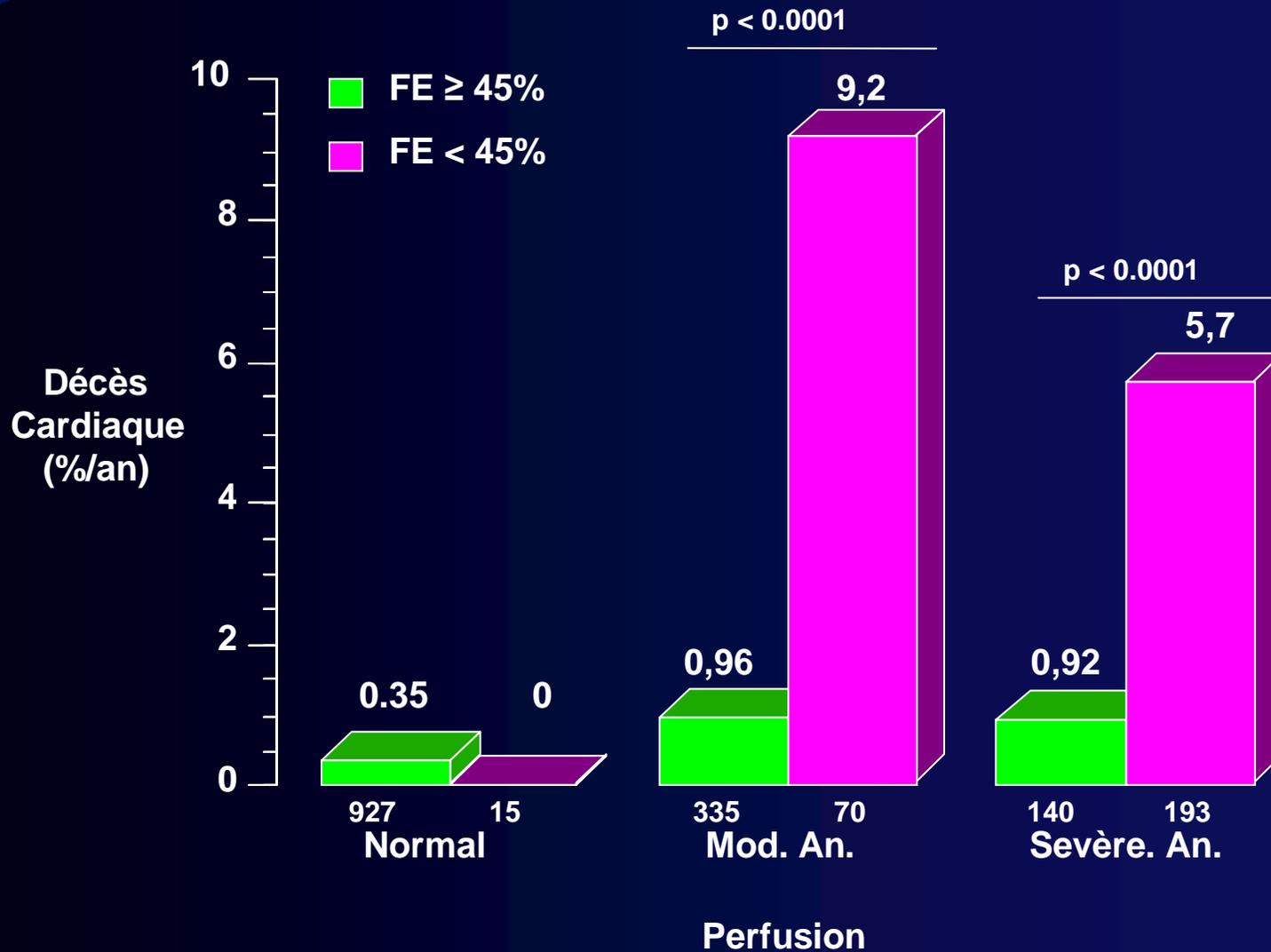
Fonction Ventriculaire Gauche



FEVG : G-SPECT



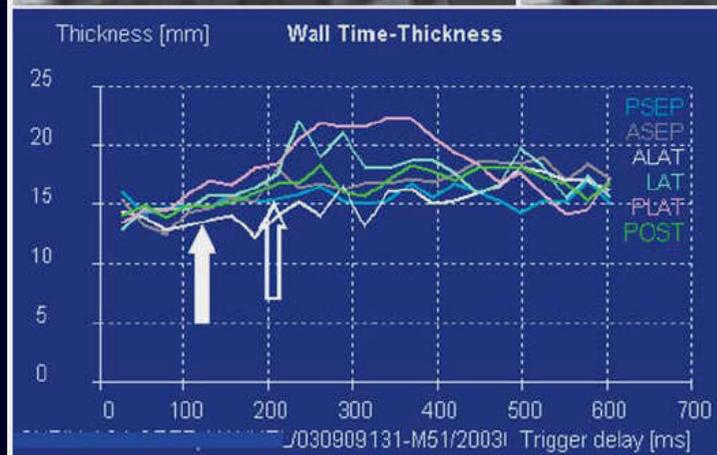
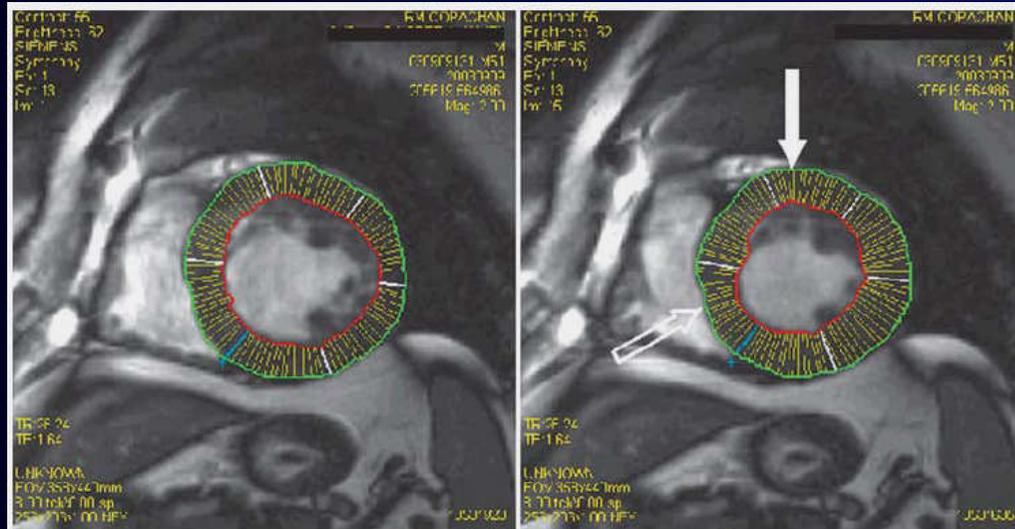
FEVG : G-SPECT



FEVG : IRM

- Etat d'équilibre
- Excellente définition endocarde / épicarde
- Absence de assumption géométrique
- Etude :
 - Des volumes TD et TS VG
 - De la FE globale
 - De la cinétique régionale
 - De l'épaississement segmentaire
- **Méthode de référence**

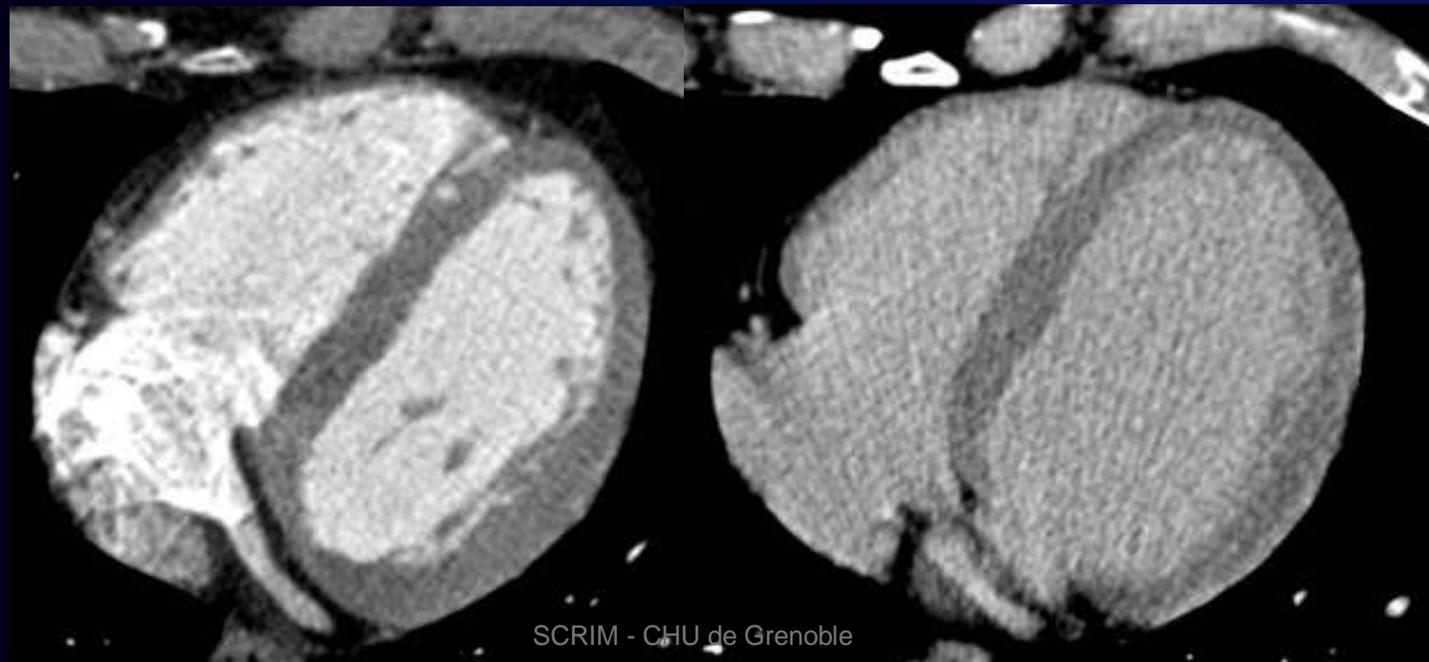
FEVG : IRM



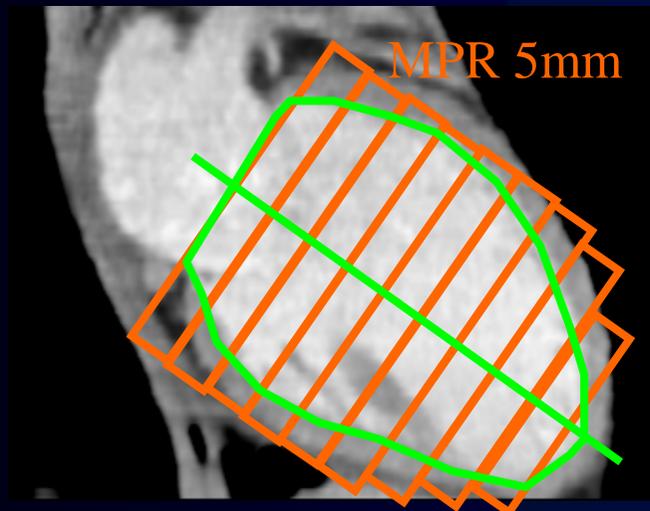
FEVG : Stress- IRM

- IRM de stress et Δc de coronaropathie :
 - 10 études
 - 654 patients (dobutamine)
 - Sensibilité 89 %
 - Spécificité 84 %

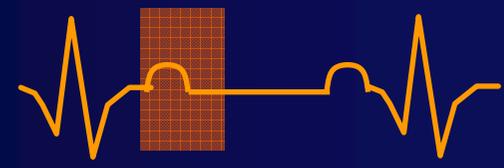
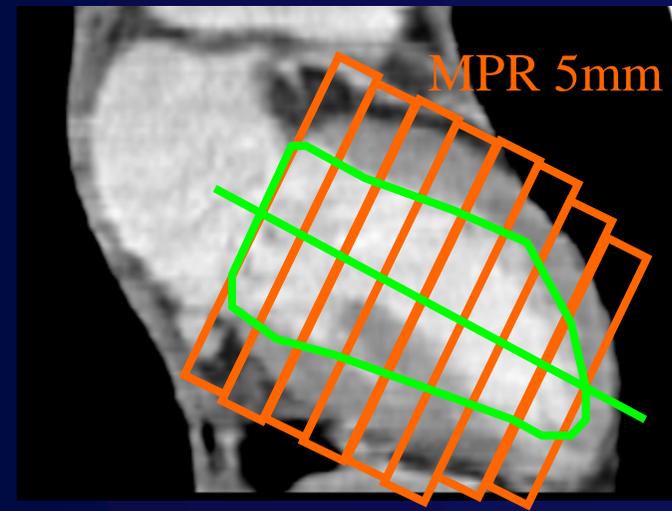
FEVG : TDM-MB



FEVG : TDM-MB

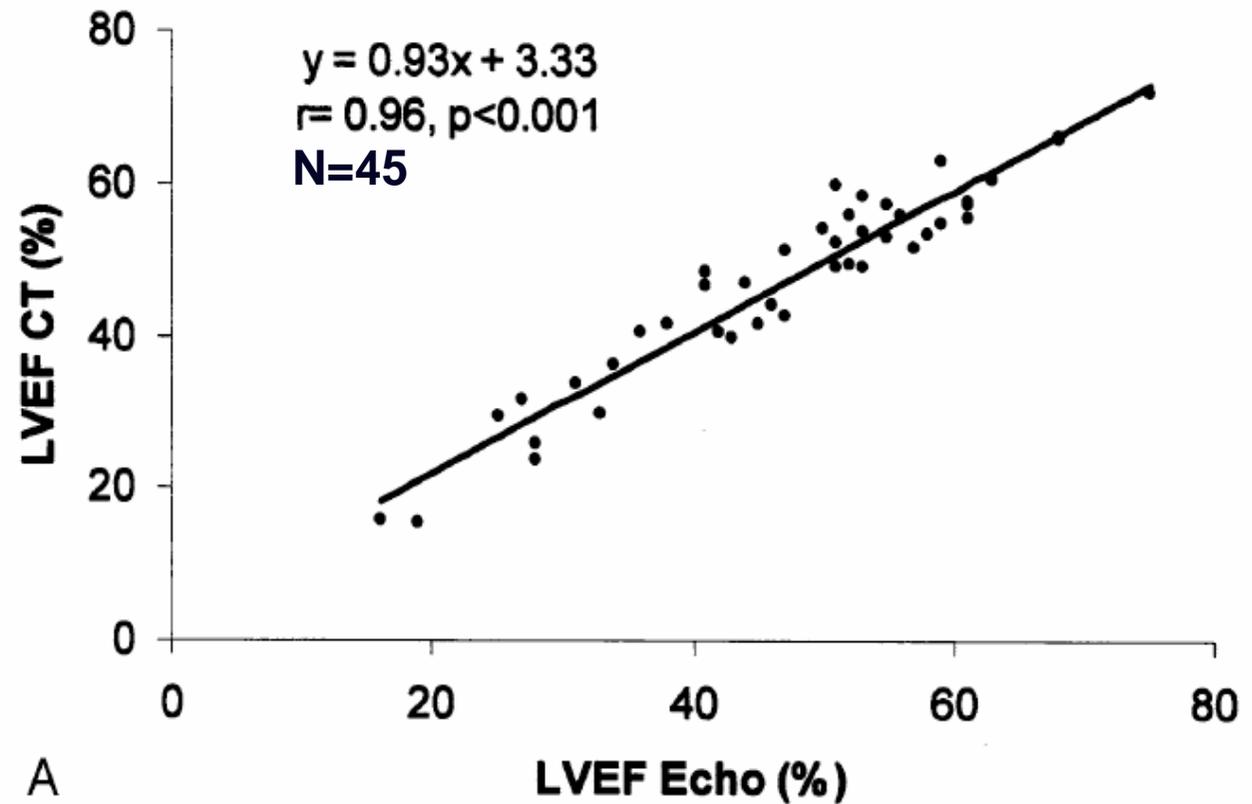
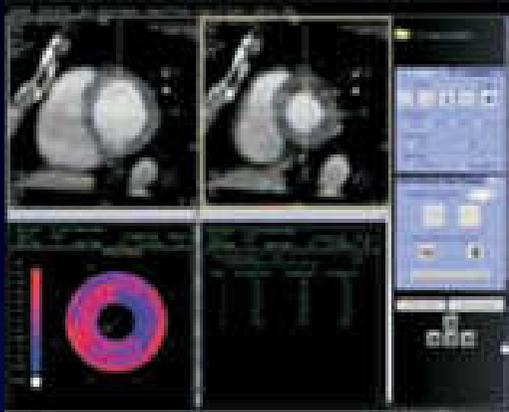


End-Diastole R-150ms

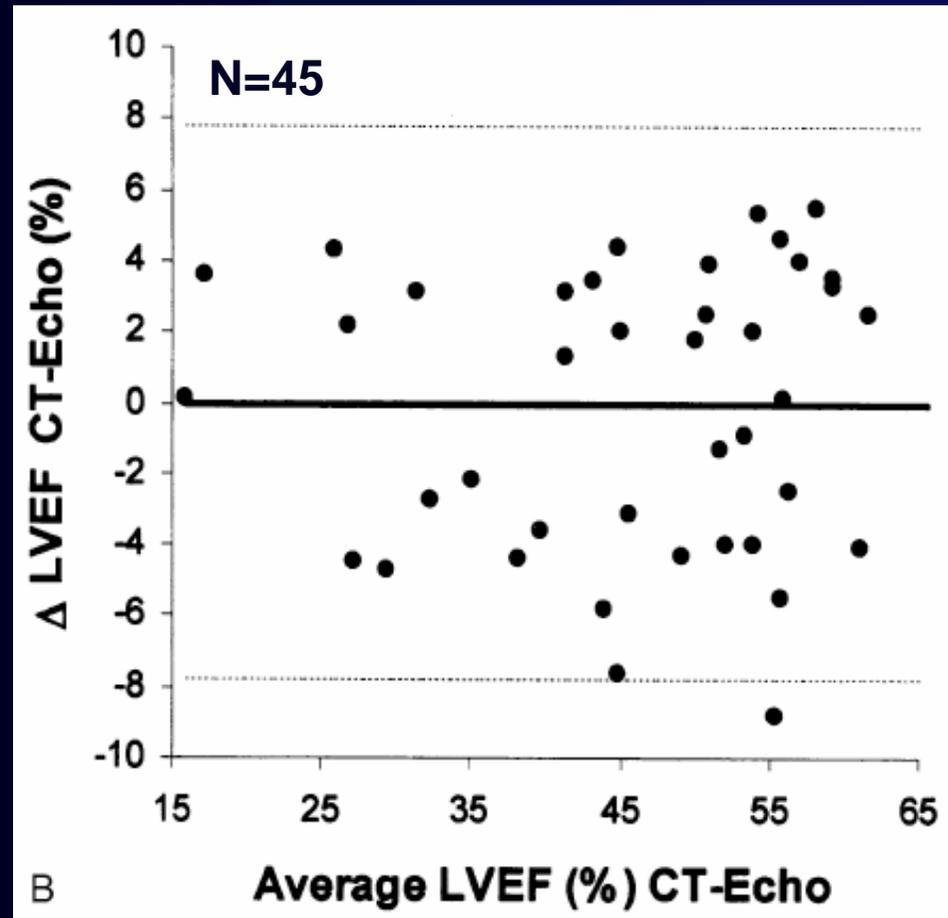


End-Systole R+150ms

FEVG TDM-MB



FEVG TDM-MB



Définition clinique

Viabilité

- Présence au sein d'un myocarde **ischémique et hypokinétique**...
- d'une quantité de cardiomyocytes " survivants" suffisante...
- pour permettre une amélioration de la **contractilité ventriculaire gauche** segmentaire et globale...
- après **revascularisation** par ACT ou PAC.

Physiopathologie

□ Myocarde hibernant (Hibernating myocardium)

Myocarde **hypocinétique et hypoperfusé (?)**

Condition ischémique

Réversibilité après revascularisation myocardique

□ Myocarde sidéré (Stunned myocardium)

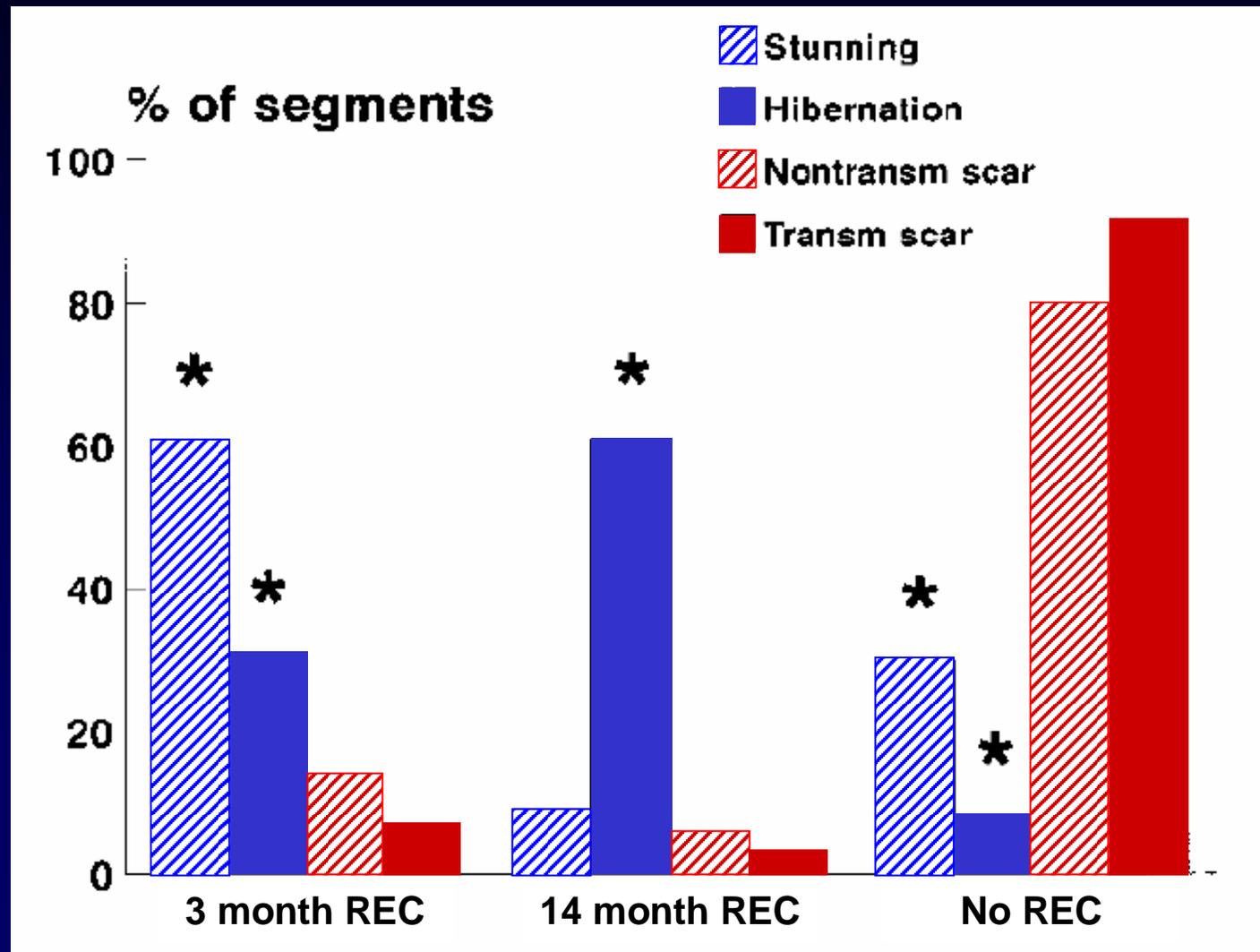
Myocarde **hypocinétique mais normoperfusé**

Conséquence d'une séquence ischémie - reperfusion

Réversibilité spontanée

□ Hibernation chronique / sidération itérative / ischémie

Délai de récupération post-opératoire



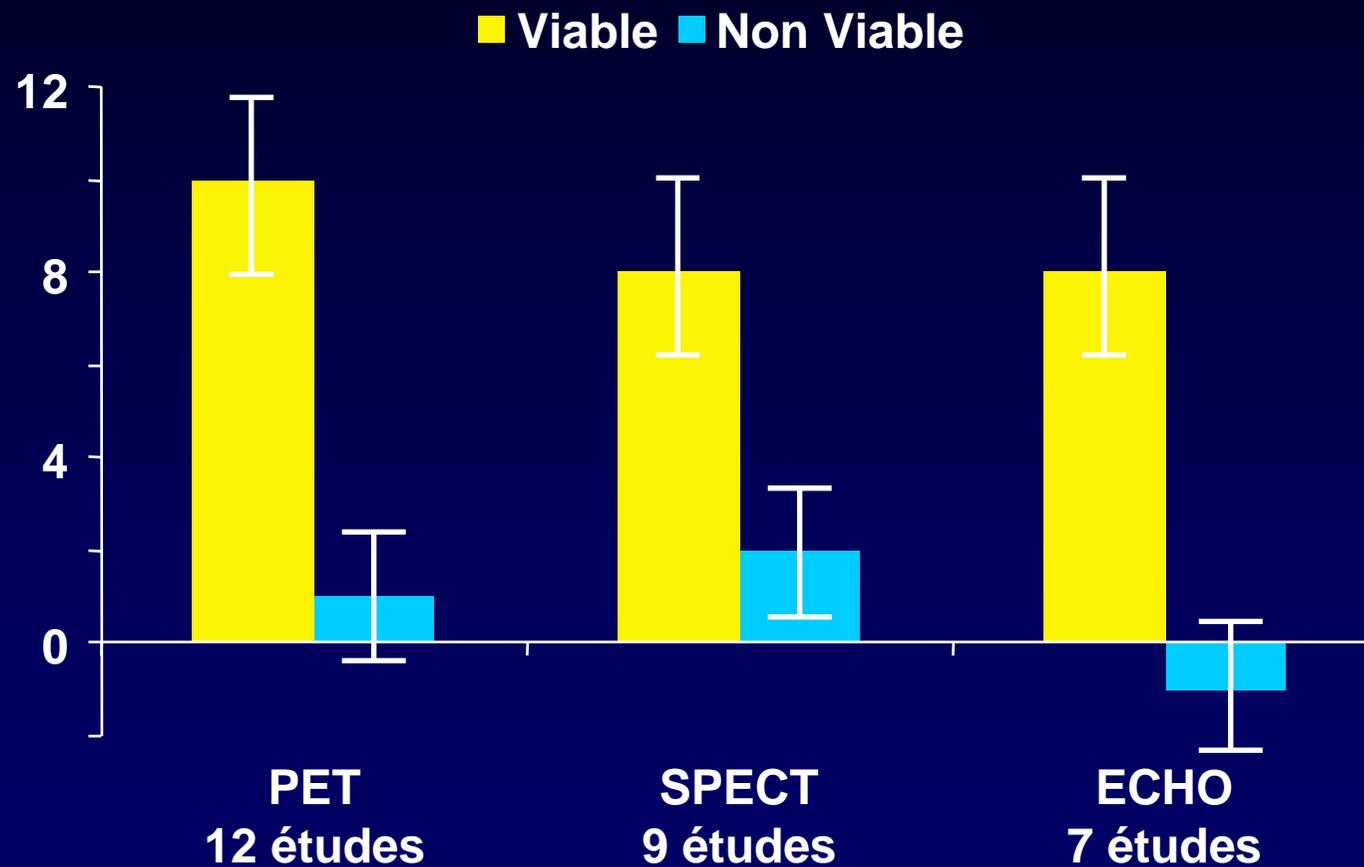
Situation clinique

Cardiopathie ischémique
Atteinte pluritronculaire
Altération de la FEVG
Insuffisance cardiaque

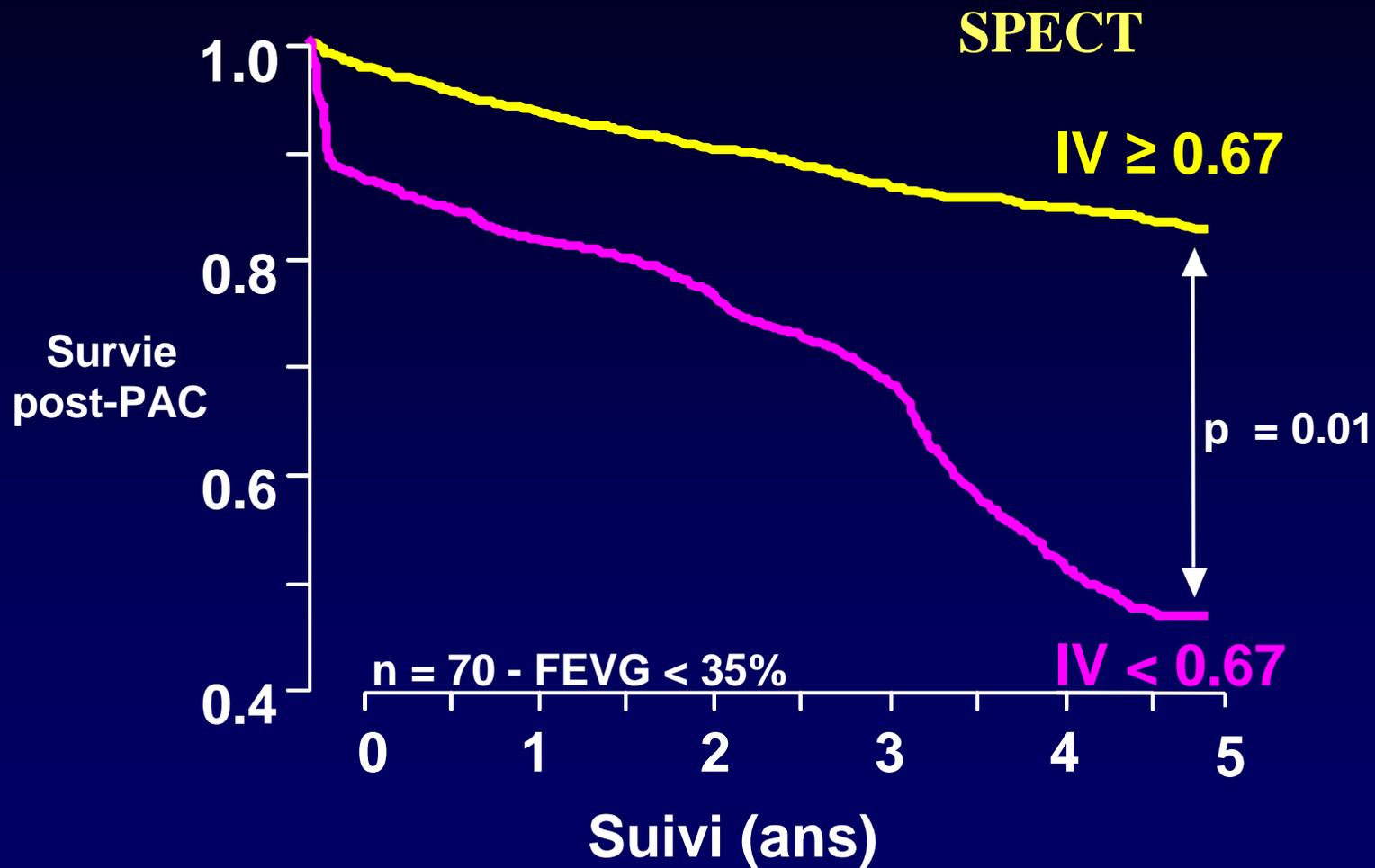
Viabilité myocardique

- Amélioration de la fonction VG après revascularisation?
- Amélioration du pronostic fonctionnel et vital?

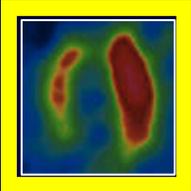
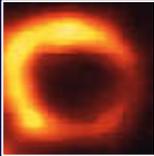
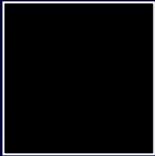
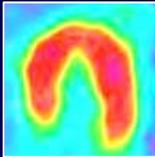
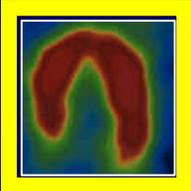
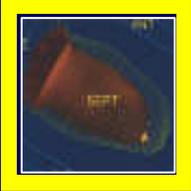
Evolution de la récupération FEVG & viabilité (1)



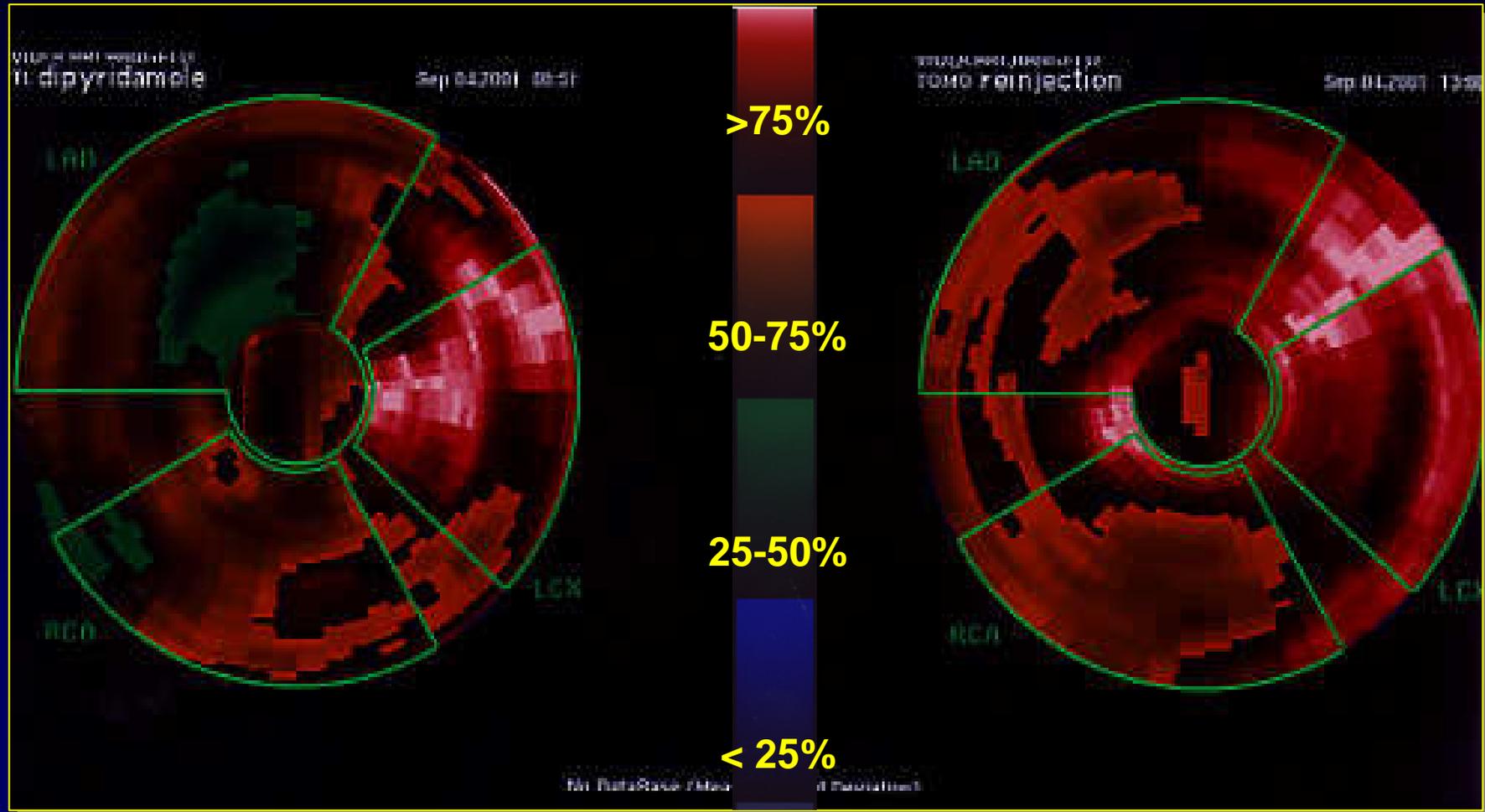
Viabilité myocardique & pronostic (2)



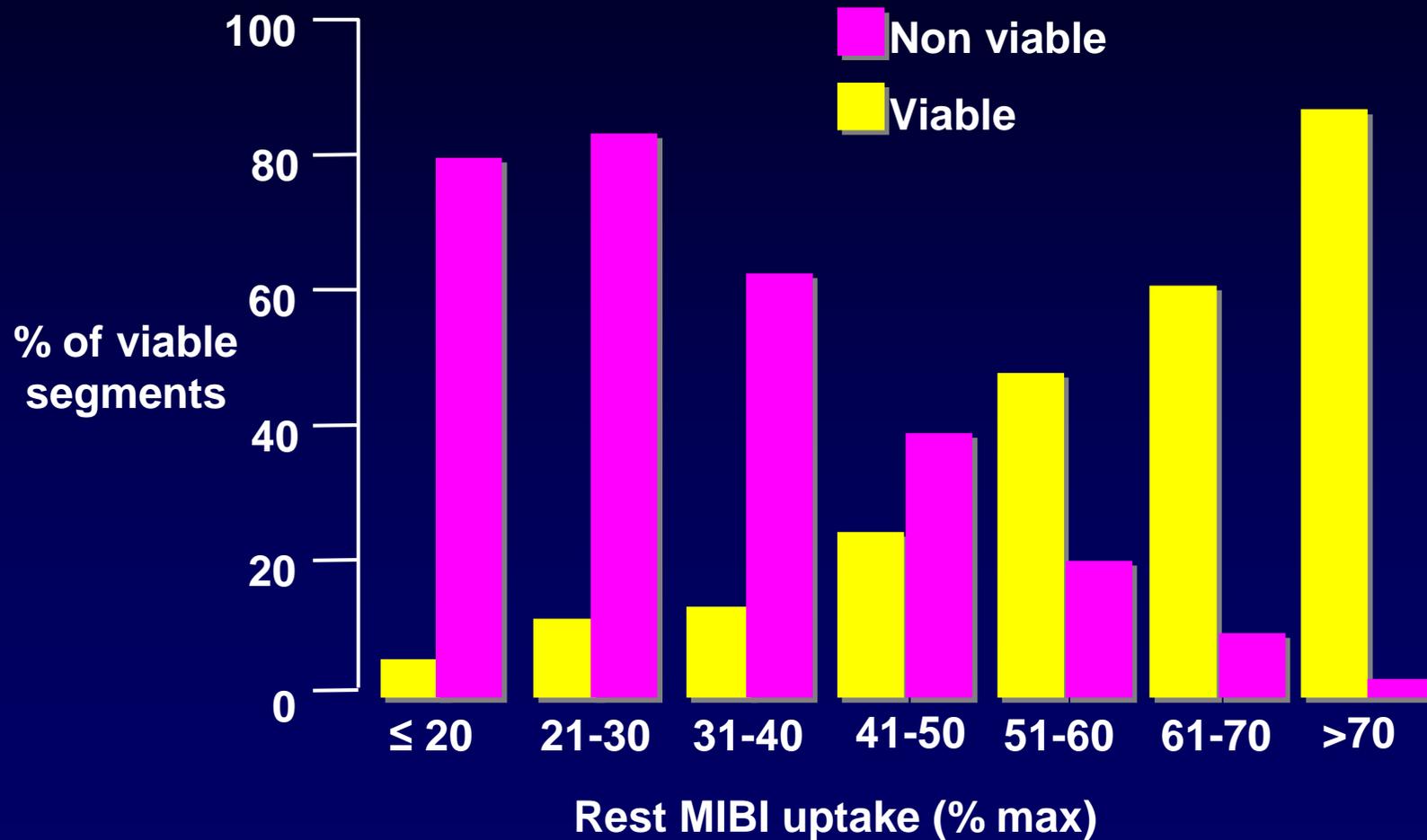
Détection de la viabilité myocardique

	SPECT	TEP	IRM	ECHO
« scar »				
Métab.				
Débit				
Fonction				

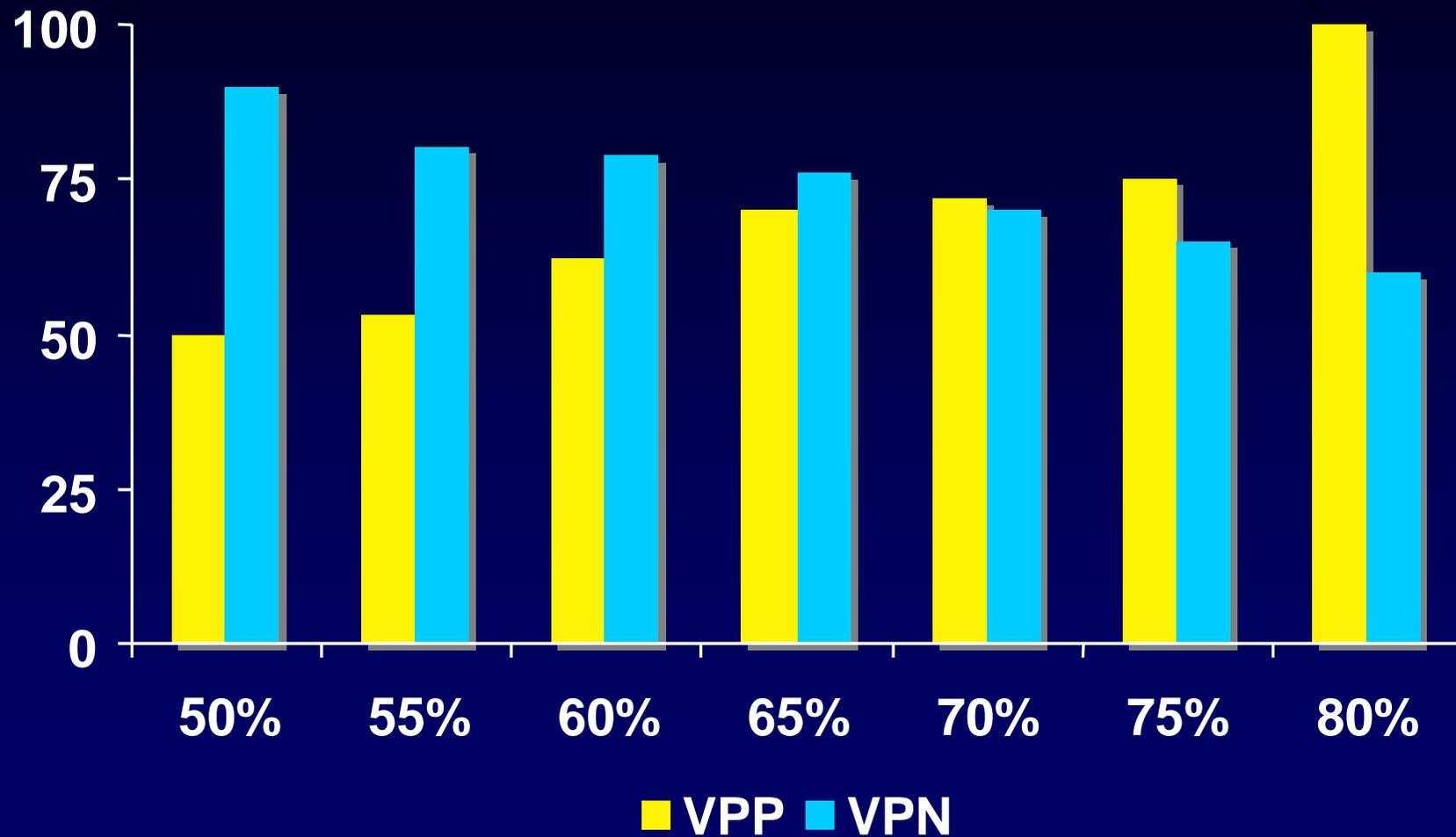
Quantification



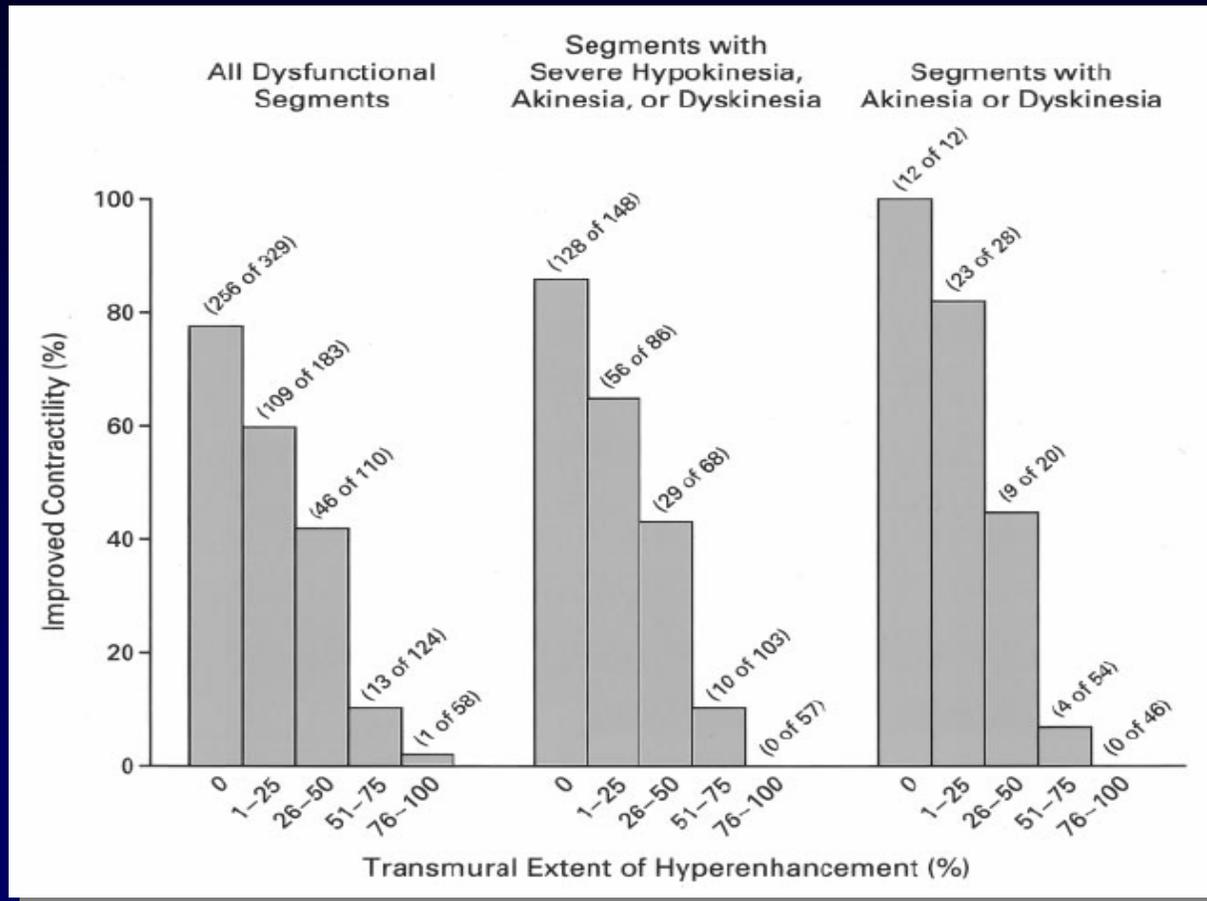
Le choix du critère diagnostique MIBI - SPECT



Le choix du critère diagnostique TI - SPECT

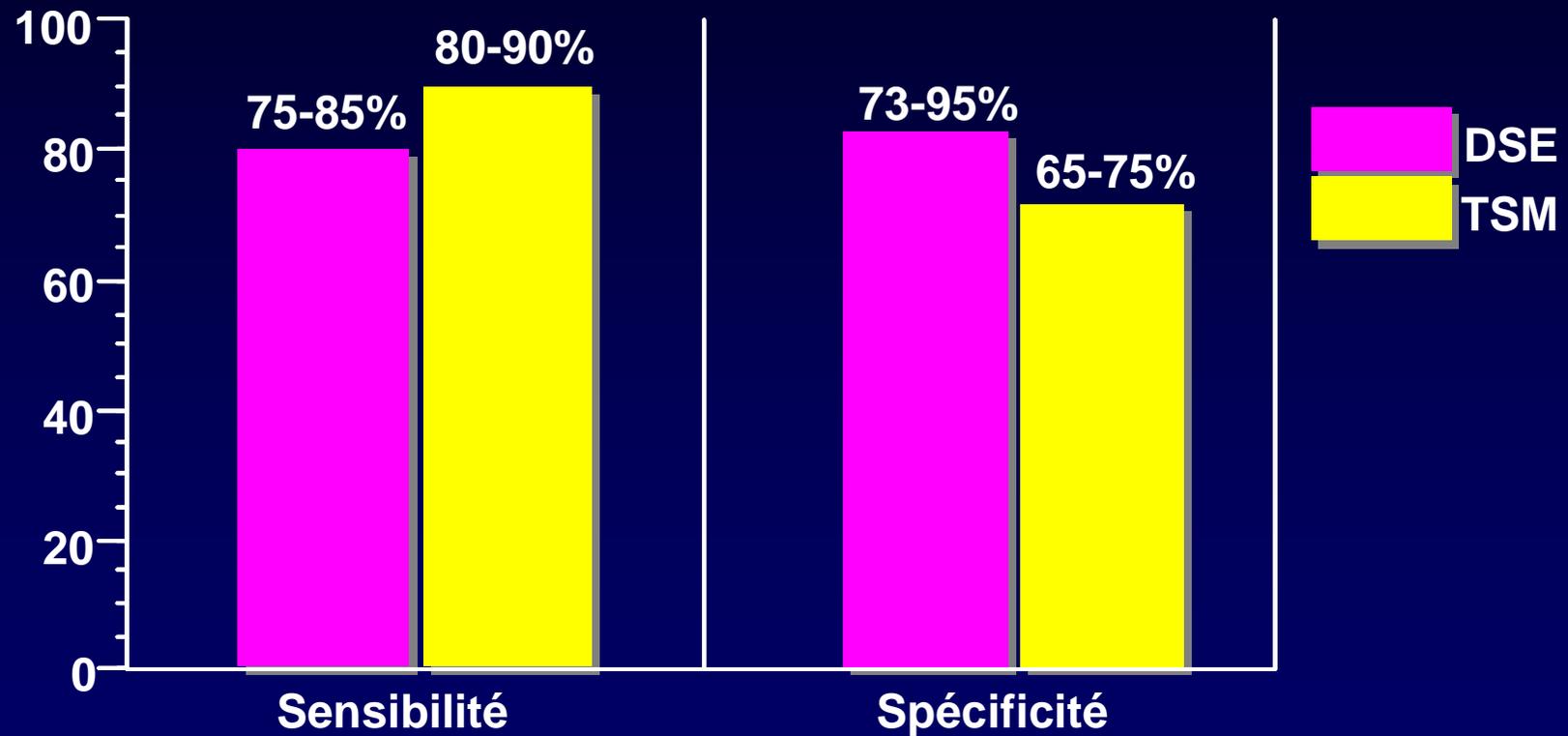


Le choix du critère diagnostique Stress-écho

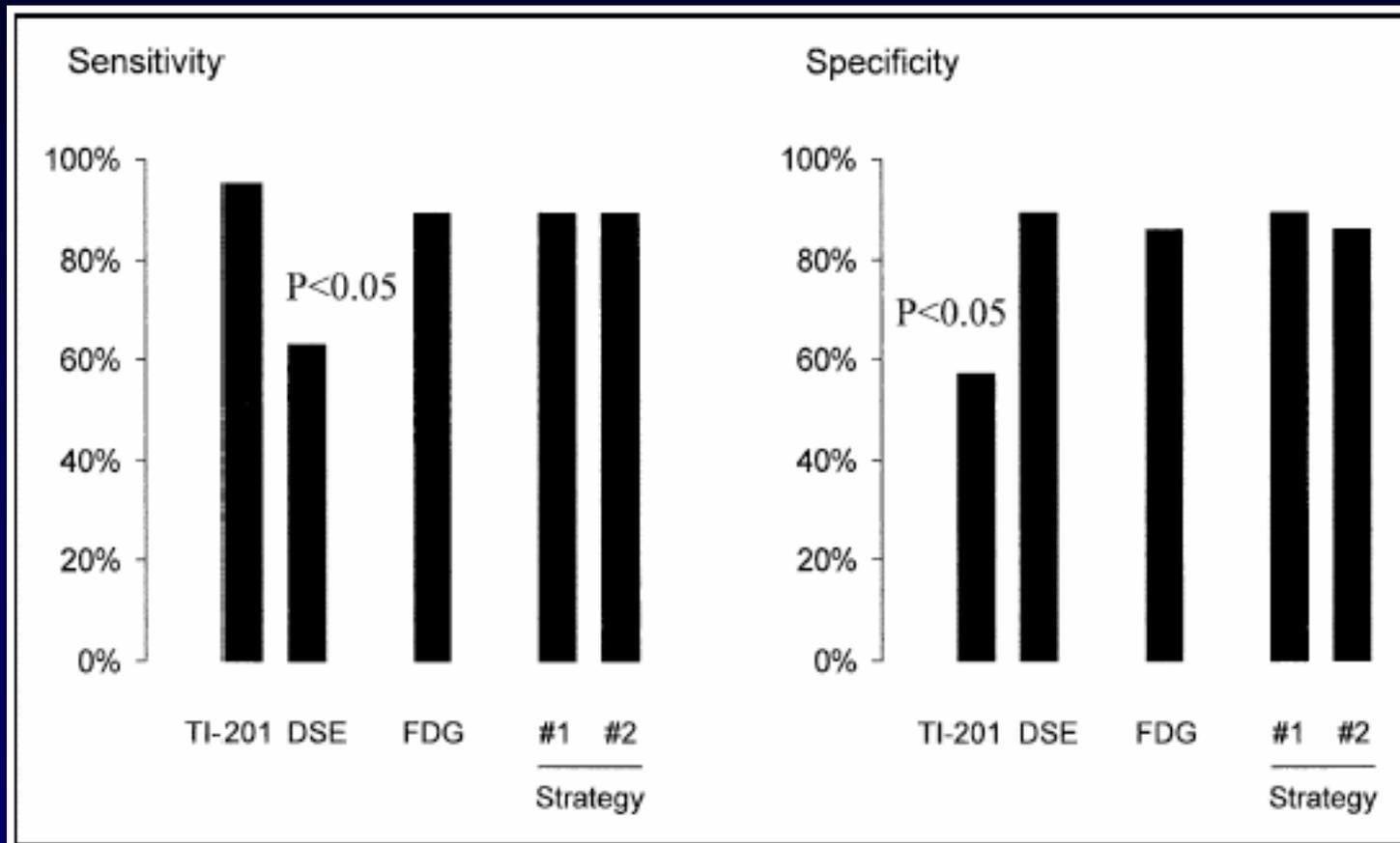


Prédiction de la récupération fonctionnelle VG

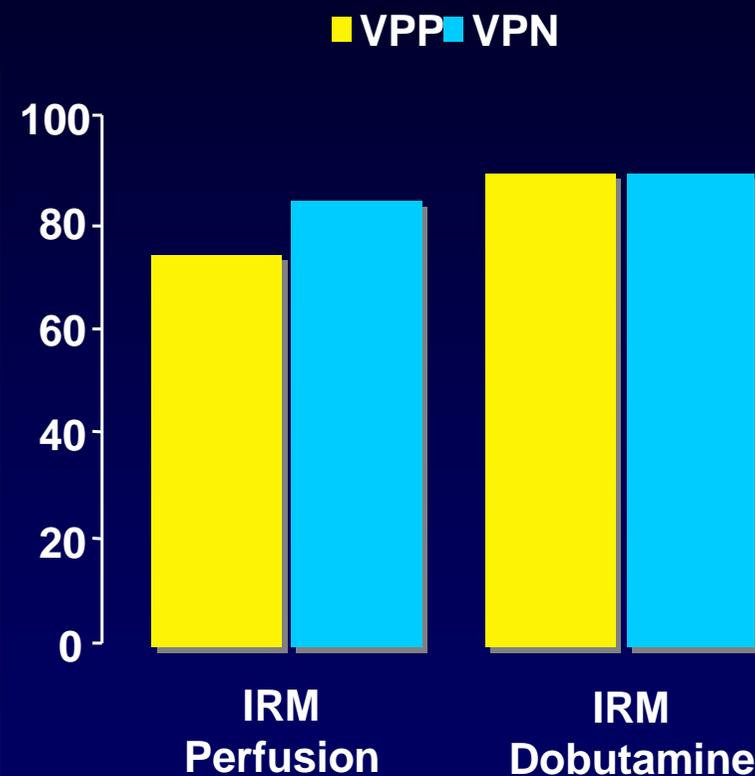
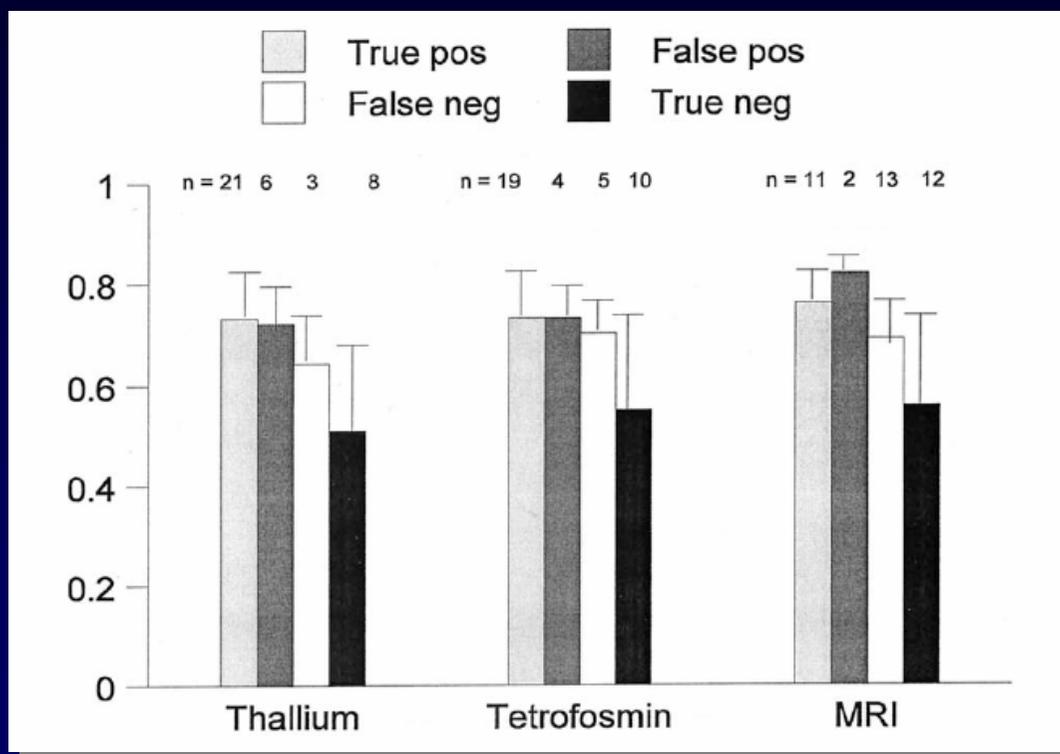
Comparaison isotopes-écho de stress



Performance des Tests Conventionnels



Performances de l'IRM



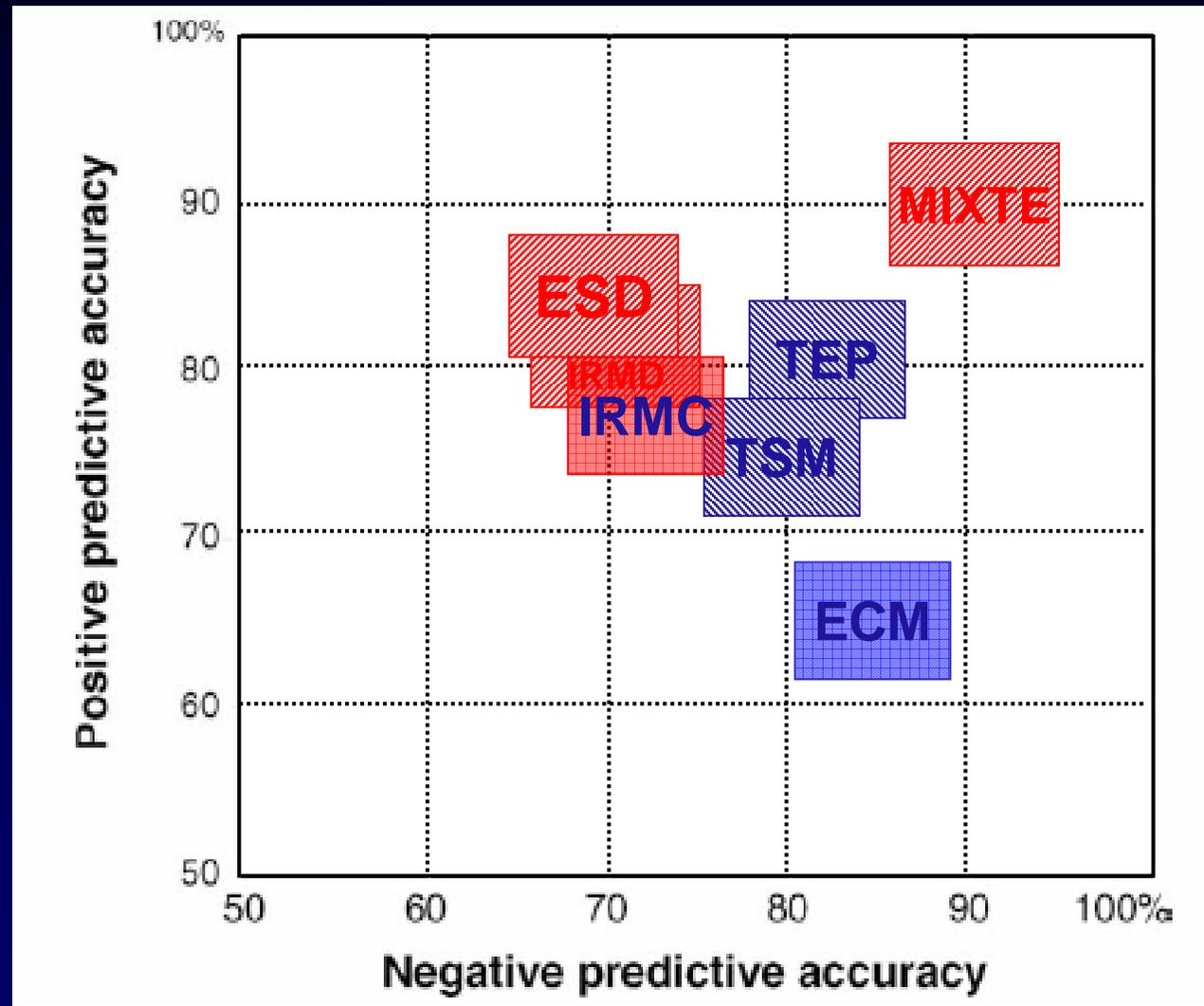
Thomson & al. *L Mag Res Imag* 2004;19:771

Shan & al. *Circulation* 2004;109:1328

Gunning & al. *JACC* 2002;39:428-35

Valeur prédictive des tests : En résumé

Baumgartner	JACC	1998
Bax	Am J Cardio	2003
Bax	Circulation	2001
Bax	JACC	1999
Bonow	Circulation	1996
Carrel	Eur J CTS	1992
Choi	Circulation	2001
Gropler J	ACC	1992
Gunning	JACC	2002
Klein	Circulation	2002
Kloner	Circulation	1998
Lucignani	Eur JNM	1992
Maddahi	Circulation	1994
Maddahi	J Nucl Med	1994
Meluzin	JACC	1998
Meza	Circulation	1997
Ragosta	Circulation	1993
Shan	Circulation	2004
Shimoni	Circulation	2003
Tamaki	Am J Cardiol	1989
Thomson	Mag Res Im	2004
Tillish	NEJHM	1986
Underwood	Eur Heart J	2004



Recommandations : Classe I ACC-AHA-ESC et viabilité

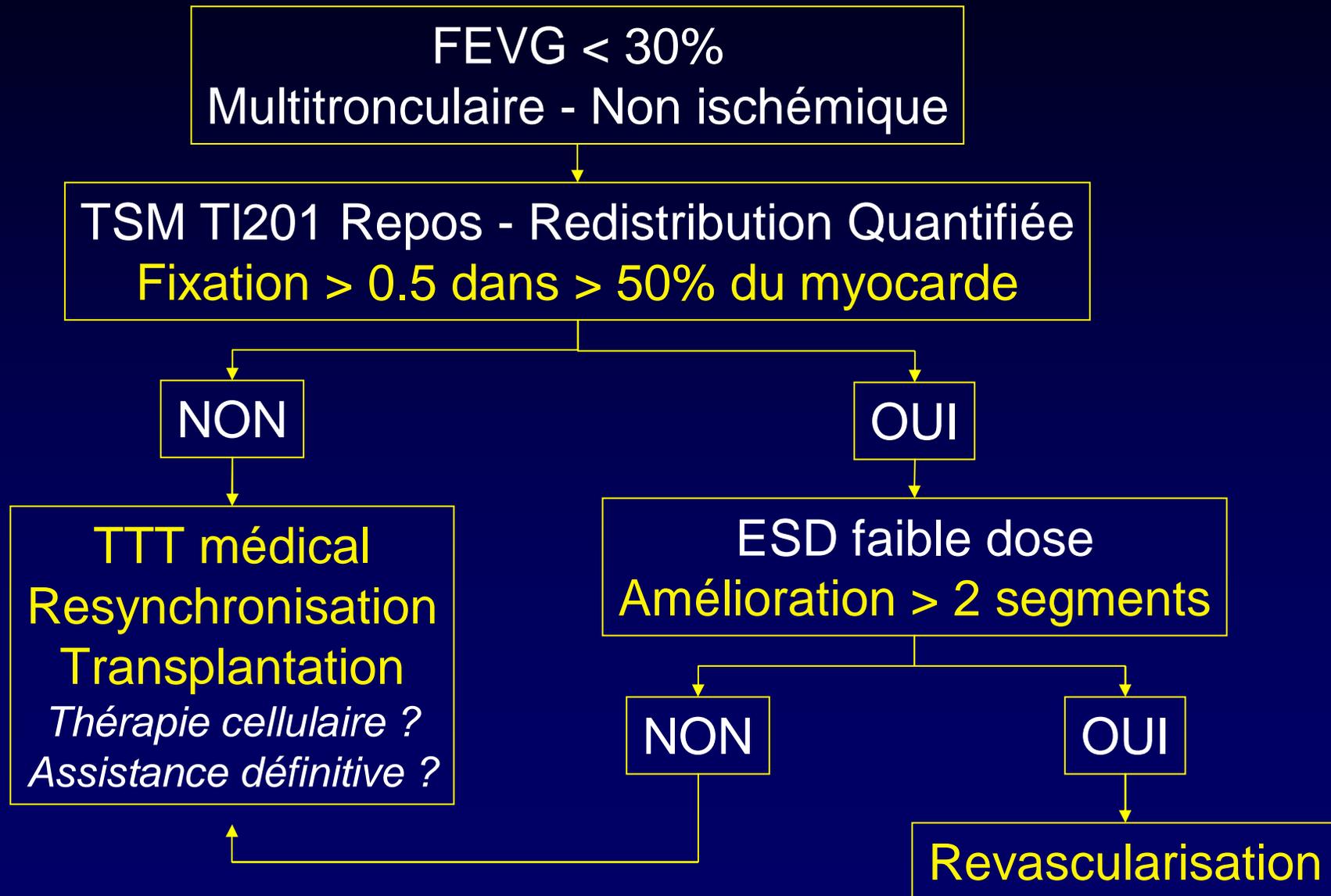
- ^{18}F FDG PET
- ^{201}Tl SPECT
- $^{99\text{m}}\text{Tc}$ -MIBI SPECT
- Stress Echo
- Stress MRI
- Contrast MRI

□ **La performance globale des tests de viabilité est entre 75 et 80%**

- ◆ Certains privilégient la VPN (TSM)
- ◆ D'autres privilégient la VPP (ESD)
- ◆ L'association des 2 offrent les meilleurs performances (proche de 90%)

□ La probabilité de récupération semble très **faible en cas de remodelage VG majeur (VTS > 130 - 150 ml)**

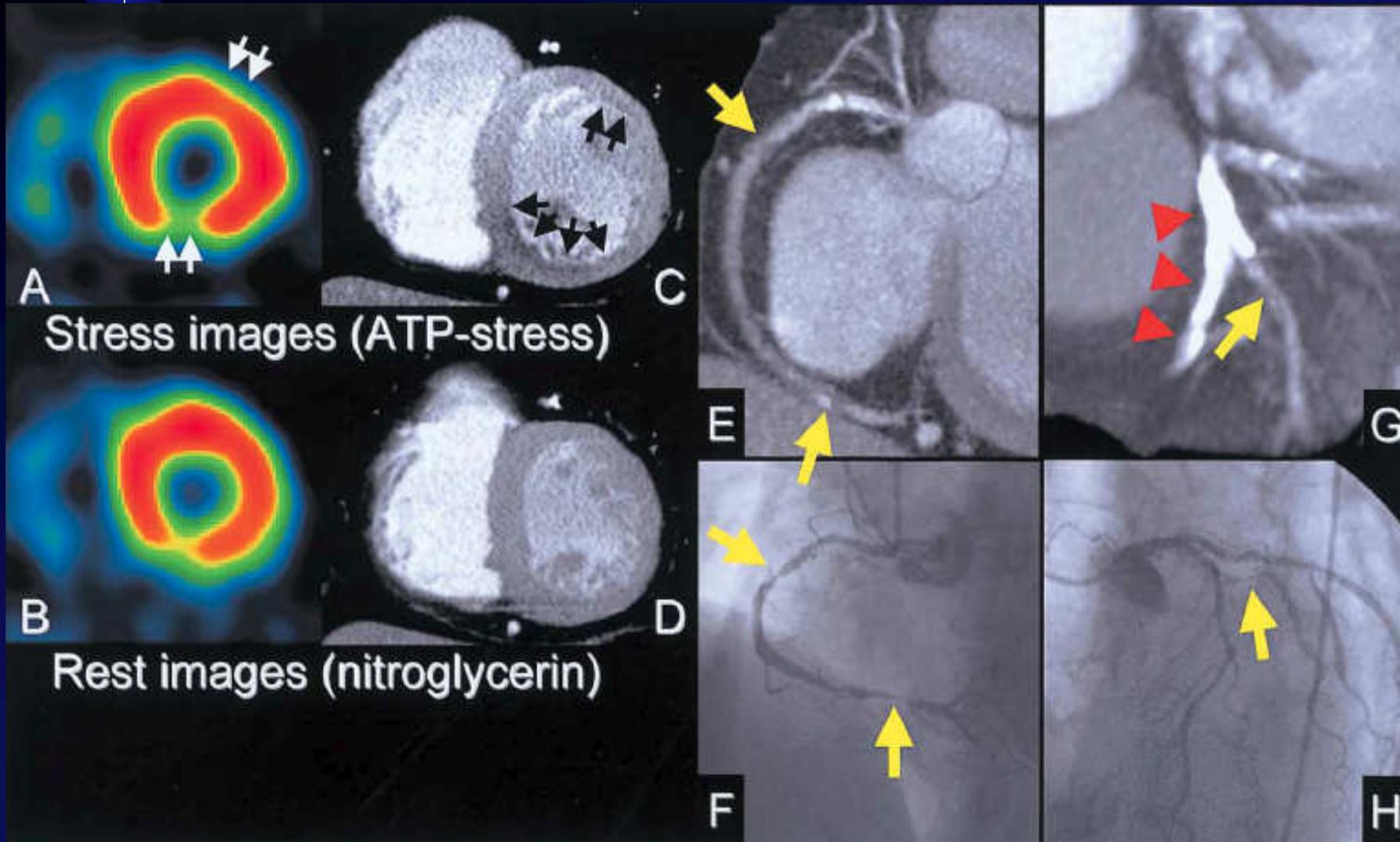
Proposition de stratégie simplifiée



L'idéal du cardiologue

- Examen largement disponible et peu coûteux
- Non invasif et peu (pas ?) irradiant pour le patient et le cardiologue
- Apportant toutes les informations nécessaires à la prise de décision thérapeutique
 - Anatomie coronaire
 - Ischémie myocardique
 - Viabilité myocardique
 - Fonction ventriculaire gauche
 - Anatomie hémodynamique intra-cardiaque
 - Comorbidité & diagnostic différentiel ?

“All heart MSCT”



Les différentes techniques en résumé

	CORO	ESD	TSM	TEP	TDM-MCB	IRM
Disponibilité (jours)	2	15	30	6	7	5
Non invasif	-	+	+	+	+	+
Irradiation (mSv)	3-8	0	15 - 30		10	0
Temps médecin (min)	20	45	10	15	30	30
Coût (€)			450	950	125	250
Coronaires	++	-	-	-	++	+
Plaque instable	-	-	?	?	++	?
Ischémie	-	+++	+++	++	+	++
Viabilité	-	++	++	+++	?	++
Fonction VG	++	++	++	?	+	+++
Anatomie	±	+++	-	-	+++	+++
Validation diagnostique*	15737	1063	2527	490	188	1194
Validation pronostique*	3154	247	423	92	9	82

Medline hits : CAD - prognosis or diagnosis