

Place de l'ECHOGRAPHIE cardiaque dans l'Insuffisance Cardiaque Aiguë

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National des
Cardiologues des
Hôpitaux

DÉCLARATION DE RELATIONS PROFESSIONNELLES

Je n'ai pas de lien d'intérêt potentiel à déclarer

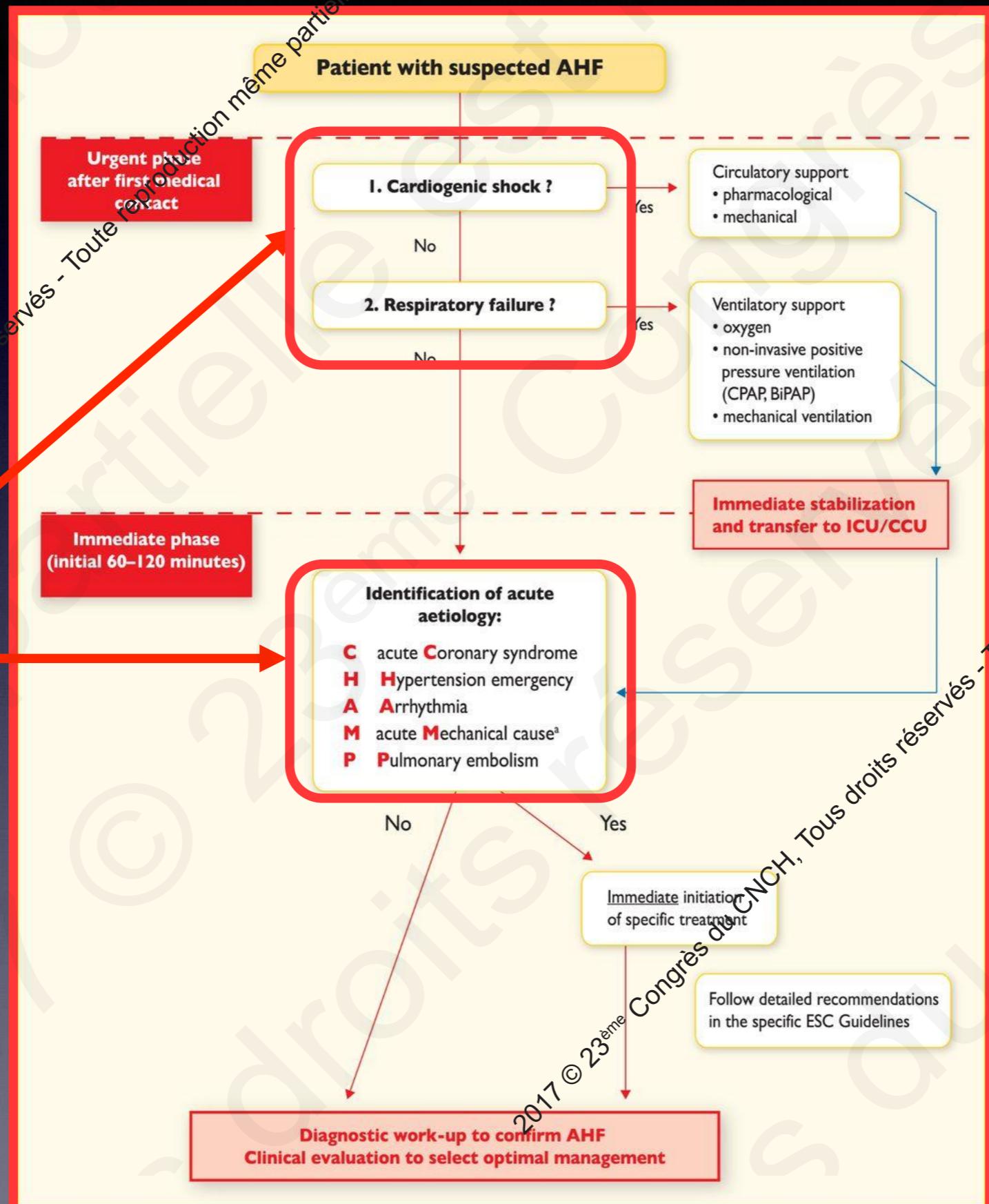
Insuffisance cardiaque

- **IC=Syndrome clinique** caractérisé par des symptômes typiques (dyspnée, OMI et fatigue) accompagnés de signes cliniques (crépitants, TJ) causé par une anomalie cardiaque structurelle ou fonctionnelle conduisant à une réduction du débit cardiaque et/ou à l'augmentation des pressions intra cardiaques au repos ou à l'effort
- **ICA:** Survenue rapide ou aggravation des symptômes d'une IC nécessitant une prise en charge rapide le plus souvent par une hospitalisation

PRISE EN CHARGE INITIALE D'UN PATIENT SUSPECT D'INSUFFISANCE CARDIAQUE AIGUE

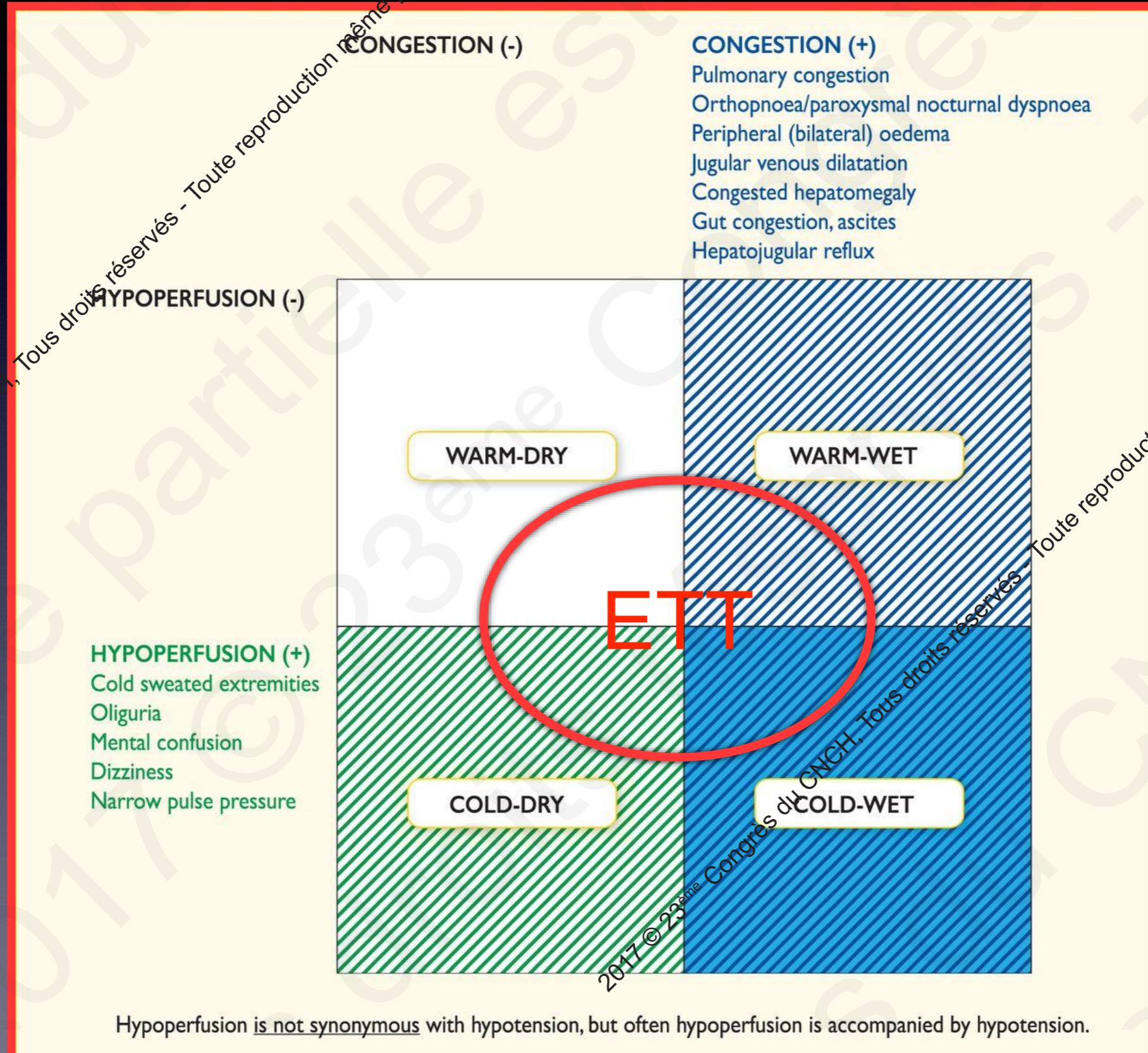
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ETT



Profil clinique des patients en ICA

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L'ECOCARDIOGRAPHIE

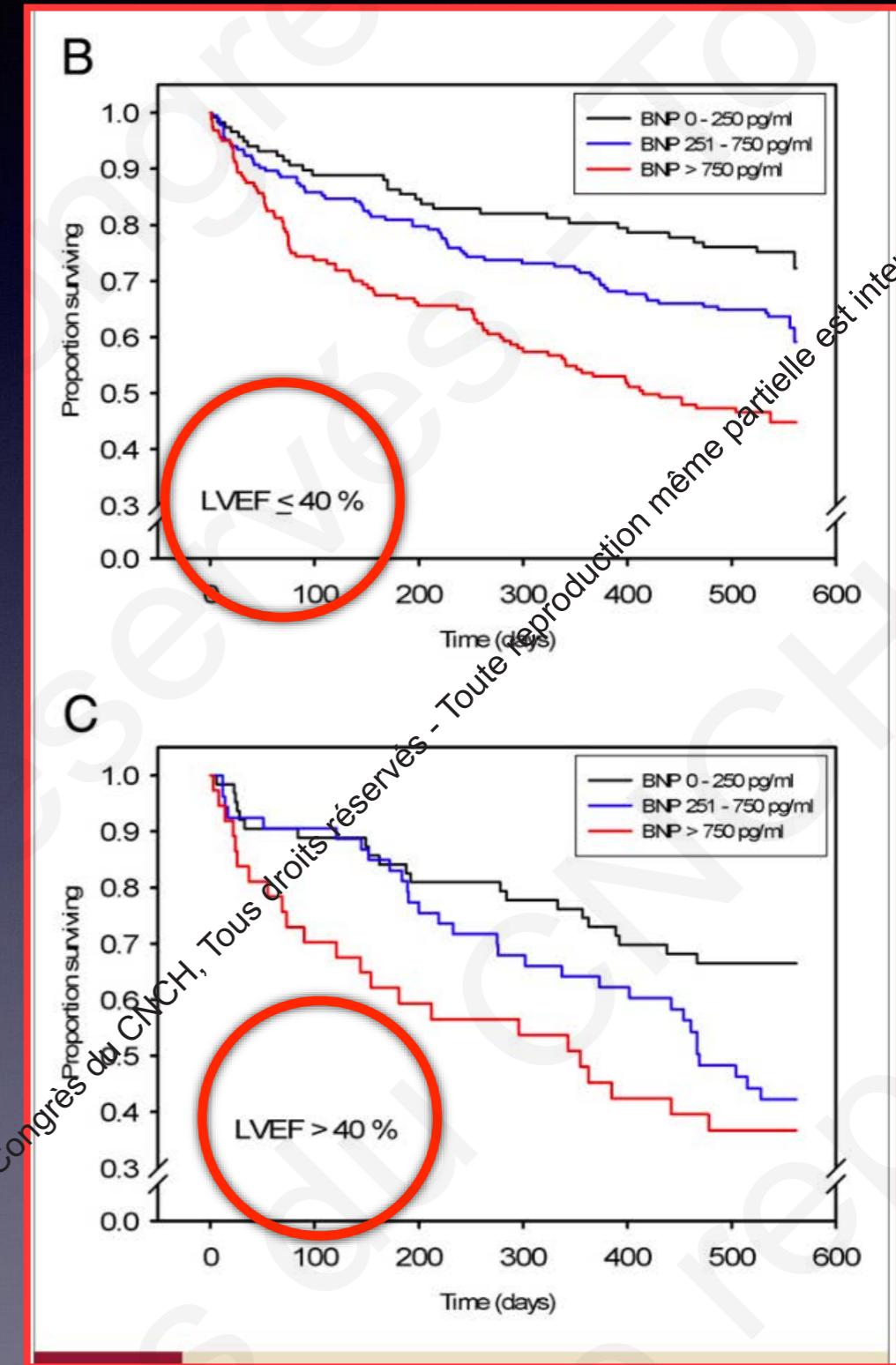
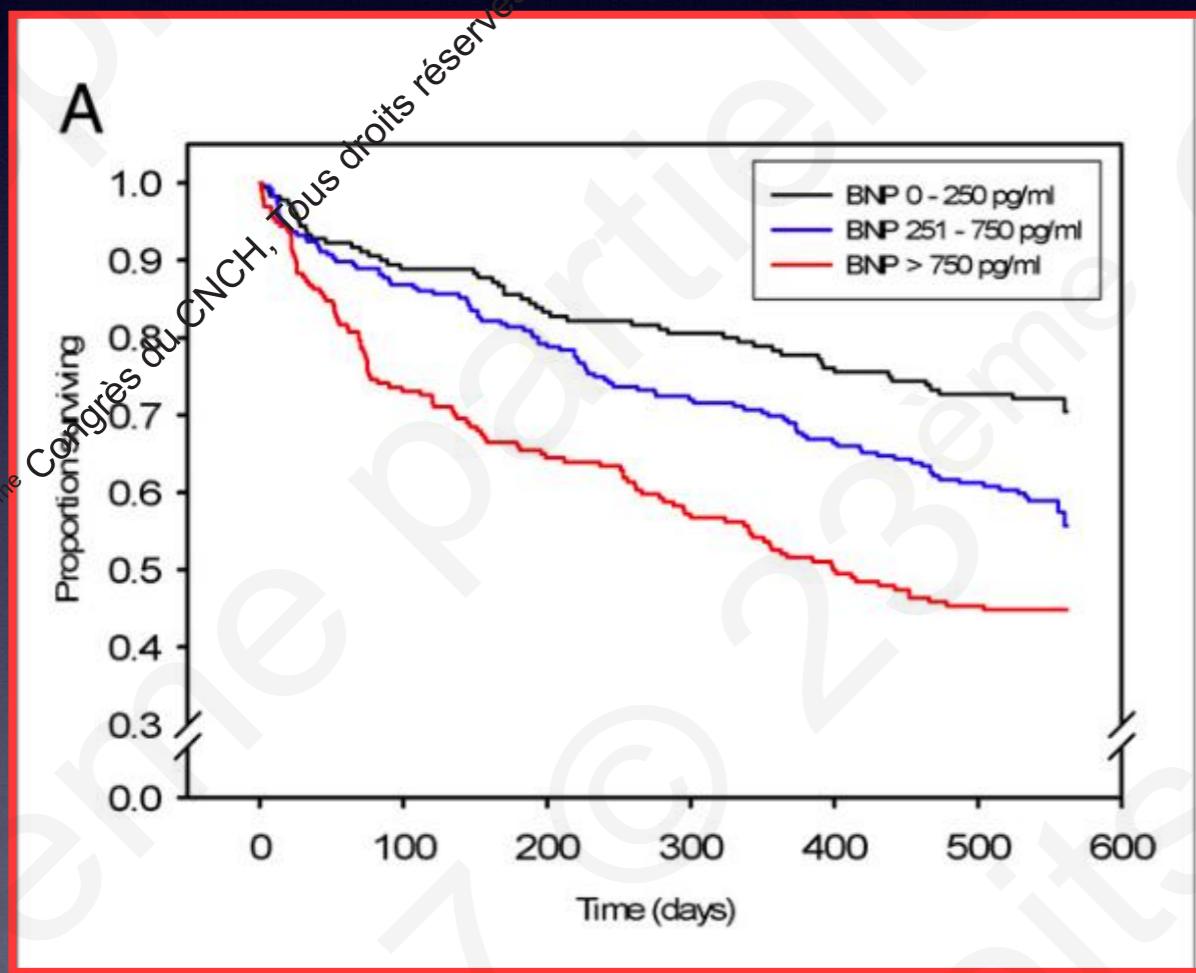
CONTRIBUE A

- ✓ Affirmer le diagnostic d'ICA, précisant l'atteinte myocardique (Systolique et/ou diastolique) , valvulaire, péricardique, endocardique
 - ✓ Trouver la ou les causes
 - ✓ Guider la thérapeutique
 - ✓ Estimer le pronostic
- ✓ Evaluation dynamique de l'ICA

CAR

- ✓ Facilité d'accès,
- ✓ Mobilité,
- ✓ Safe,
- ✓ Reproductible,
- ✓ Non Invasif

Le taux de BNP ou NT Pro-BNP ne suffit-il pas ?



Le taux de BNP ou NT Pro-BNP ne suffit-il pas ?

Table 12.3 Causes of elevated concentrations of natriuretic peptides^{522–524}

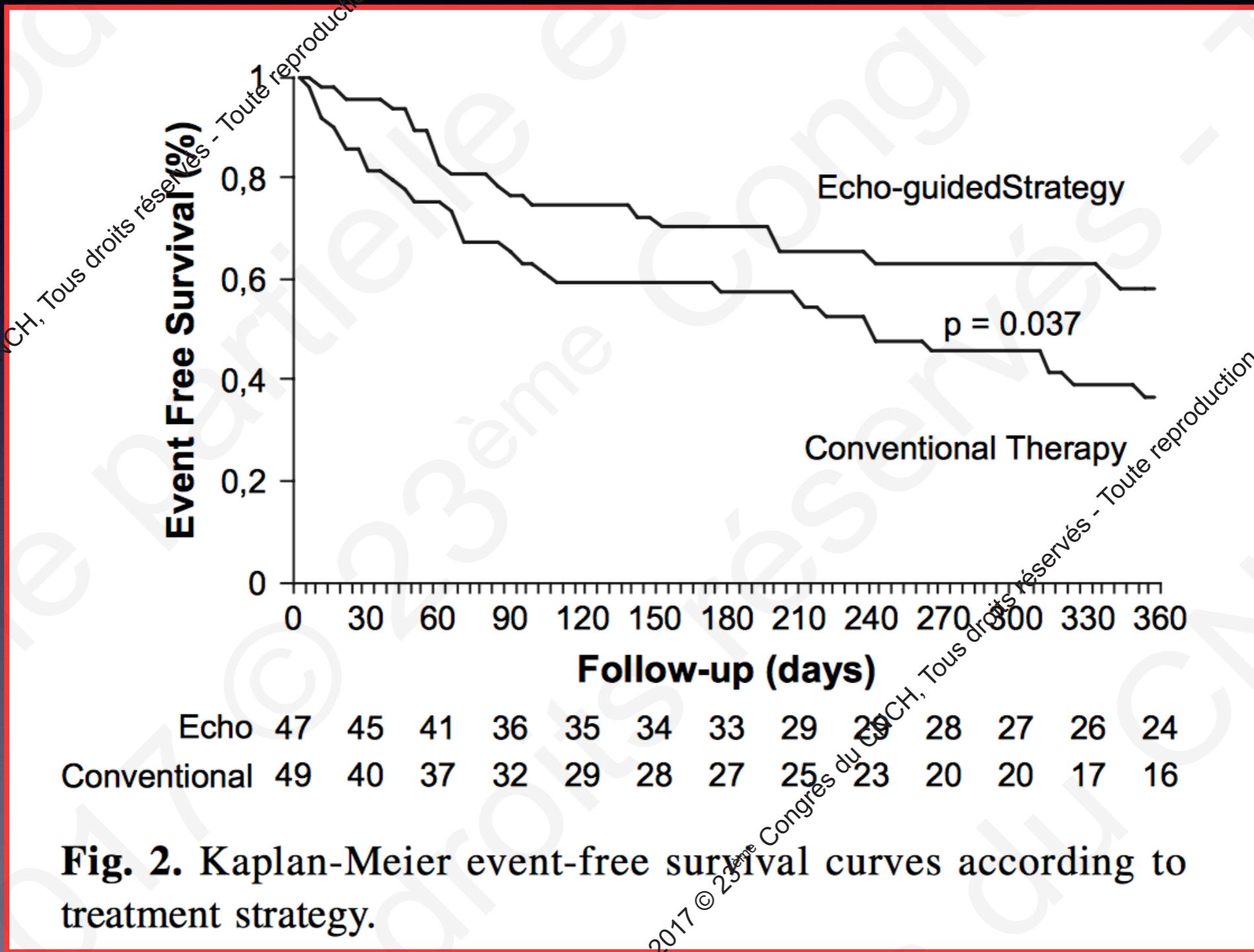
Cardiac	Heart failure Acute coronary syndromes Pulmonary embolism Myocarditis Left ventricular hypertrophy Hypertrophic or restrictive cardiomyopathy Valvular heart disease Congenital heart disease Atrial and ventricular tachyarrhythmias Heart contusion Cardioversion, ICD shock
	Surgical procedures involving the heart Pulmonary hypertension
Non-cardiac	Advanced age Ischaemic stroke Subarachnoid haemorrhage Renal dysfunction Liver dysfunction (mainly liver cirrhosis with ascites) Paraneoplastic syndrome Chronic obstructive pulmonary disease Severe infections (including pneumonia and sepsis) Severe burns Anaemia Severe metabolic and hormone abnormalities (e.g. thyrotoxicosis, diabetic ketoacidosis)

HFpEF = heart failure with preserved ejection fraction; HFrEF = heart failure with reduced ejection fraction; ICD = implantable cardioverter defibrillator.

Immediate echocardiography is mandatory only in patients with haemodynamic instability (particularly in cardiogenic shock) and in patients suspected of acute life-threatening structural or functional cardiac abnormalities (mechanical complications, acute valvular regurgitation, aortic dissection).

Early echocardiography should be considered in all patients with de novo AHF and in those with unknown cardiac function; however, the optimal timing is unknown (preferably within 48 h from admission, if the expertise is available).

Efficacité d'une stratégie écho-guidée



ETIOLOGIES DES INSUFFISANCES CARDIAQUES

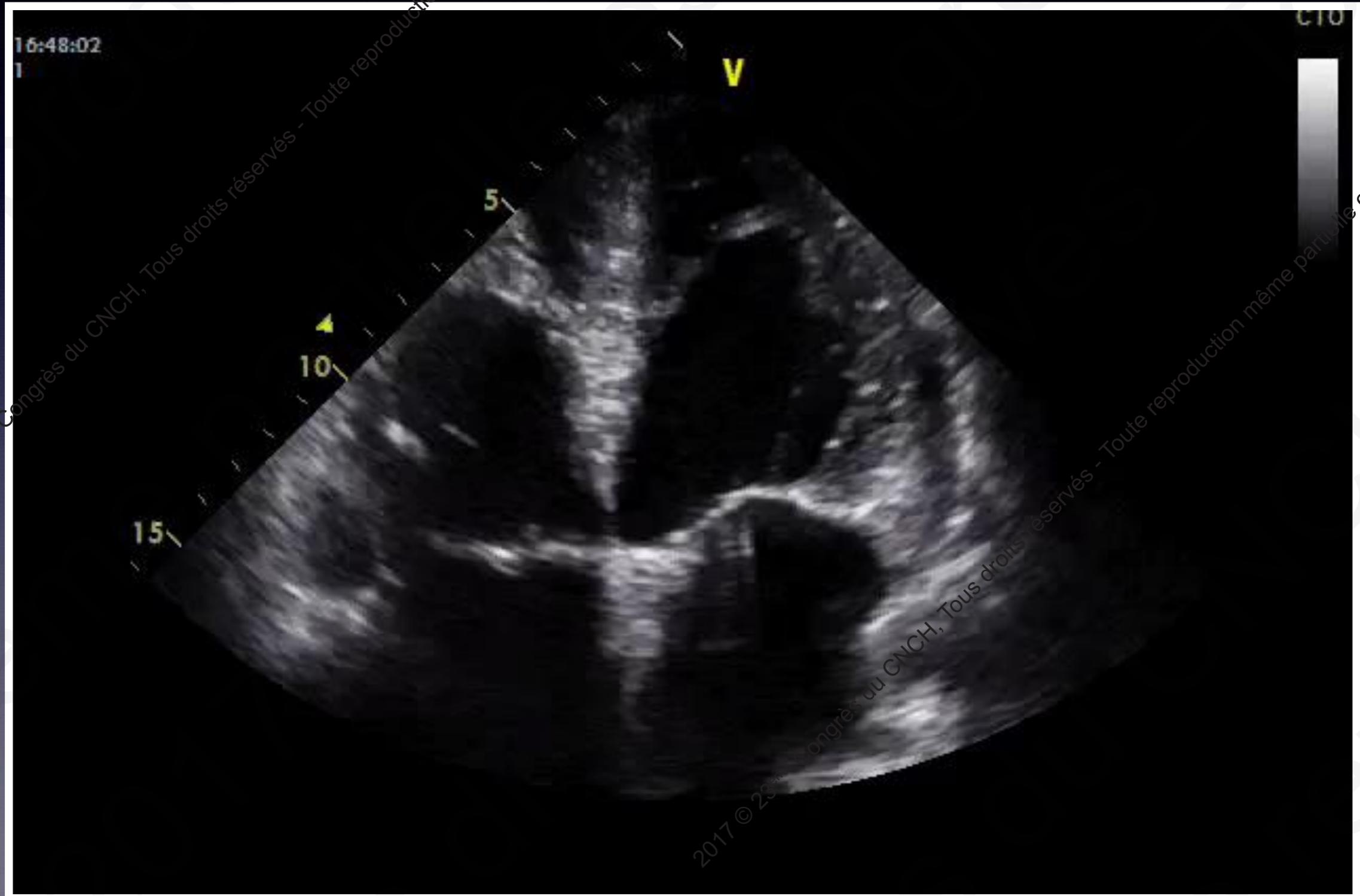
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DISEASED MYOCARDIUM		
Ischaemic heart disease	Myocardial scar	
	Myocardial stunning/hibernation	
	Epicardial coronary artery disease	
	Abnormal coronary microcirculation	
	Endothelial dysfunction	
Toxic damage	Recreational substance abuse	Alcohol, cocaine, amphetamine, anabolic steroids.
	Heavy metals	Copper, iron, lead, cobalt.
	Medications	Cytostatic drugs (e.g. anthracyclines), immunomodulating drugs (e.g. interferons monoclonal antibodies such as trastuzumab, cetuximab), antidepressant drugs, antiarrhythmics, non-steroidal anti-Inflammatory drugs, anaesthetics.
	Radiation	
Immune-mediated and inflammatory damage	Related to infection	Bacteria, spirochaetes, fungi, protozoa, parasites (Chagas disease), rickettsiae, viruses (HIV/AIDS).
	Not related to infection	Lymphocytic/giant cell myocarditis, autoimmune diseases (e.g. Graves' disease, rheumatoid arthritis, connective tissue disorders, mainly systemic lupus erythematosus), hypersensitivity and eosinophilic myocarditis (Churg–Strauss).
Infiltration	Related to malignancy	Direct infiltrations and metastases.
	Not related to malignancy	Amyloidosis, sarcoidosis, haemochromatosis (iron), glycogen storage diseases (e.g. Pompe disease), lysosomal storage diseases (e.g. Fabry disease).
Metabolic derangements	Hormonal	Thyroid diseases, parathyroid diseases, acromegaly, GH deficiency, hypercortisolism, Conn's disease, Addison disease, diabetes, metabolic syndrome, phaeochromocytoma, pathologies related to pregnancy and peripartum.
	Nutritional	Deficiencies in thiamine, L-carnitine, selenium, iron, phosphates, calcium, complex malnutrition (e.g. malignancy, AIDS, anorexia nervosa), obesity.
Genetic abnormalities	Diverse forms	HCM, DCM, LV non-compaction, ARVC, restrictive cardiomyopathy (for details see respective expert documents), muscular dystrophies and laminopathies.
ABNORMAL LOADING CONDITIONS		
Hypertension		
Valve and myocardium structural defects	Acquired	Mitral, aortic, tricuspid and pulmonary valve diseases.
	Congenital	Atrial and ventricular septum defects and others (for details see a respective expert document).
Pericardial and endomyocardial pathologies	Pericardial	Constrictive pericarditis Pericardial effusion
	Endomyocardial	HES, EMF, endocardial fibroelastosis.
High output states		Severe anaemia, sepsis, thyrotoxicosis, Paget's disease, arteriovenous fistula, pregnancy.
Volume overload		Renal failure, iatrogenic fluid overload.
ARRHYTHMIAS		
Tachyarrhythmias		Atrial, ventricular arrhythmias.
Bradyarrhythmias		Sinus node dysfunctions, conduction disorders.

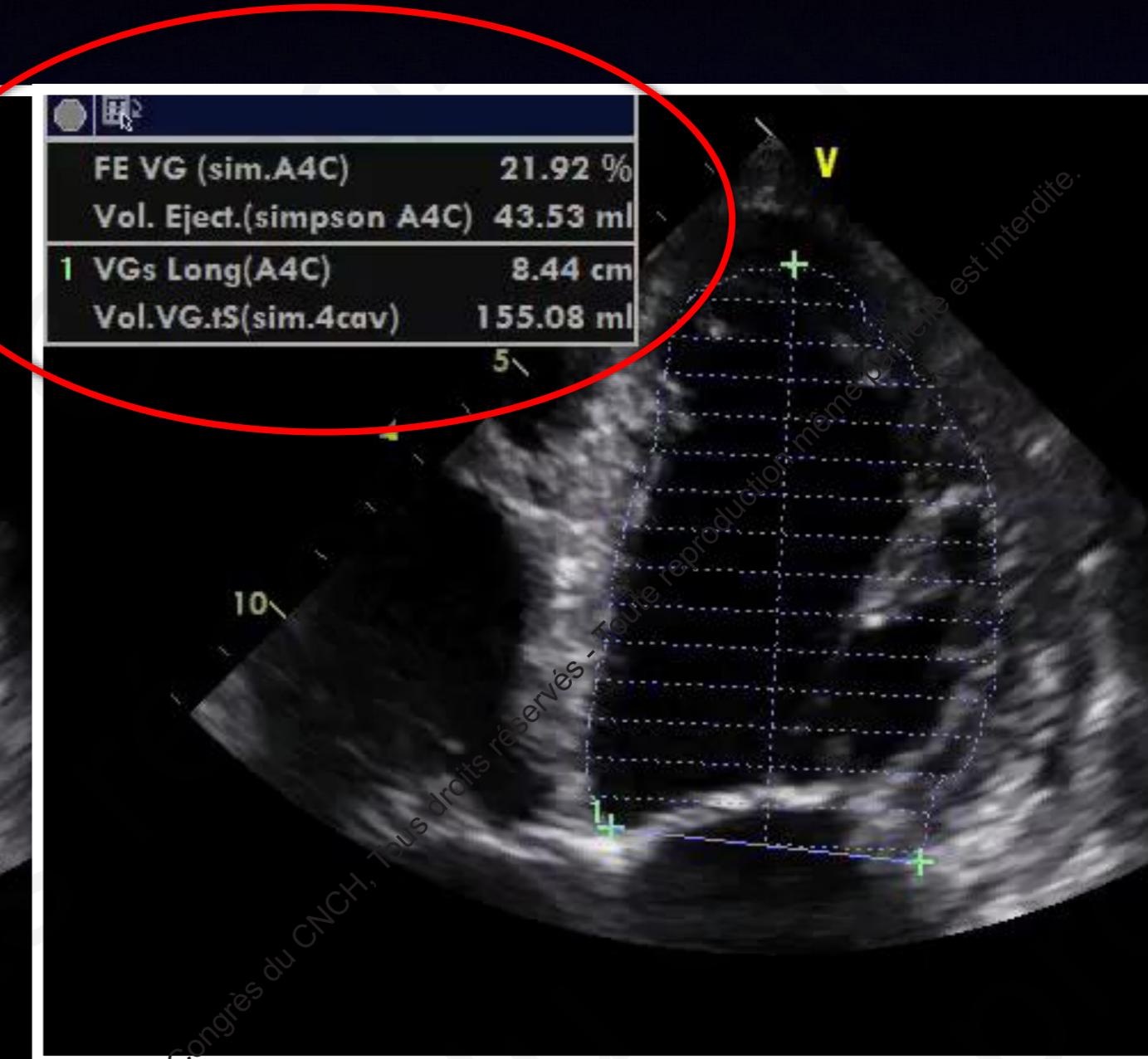
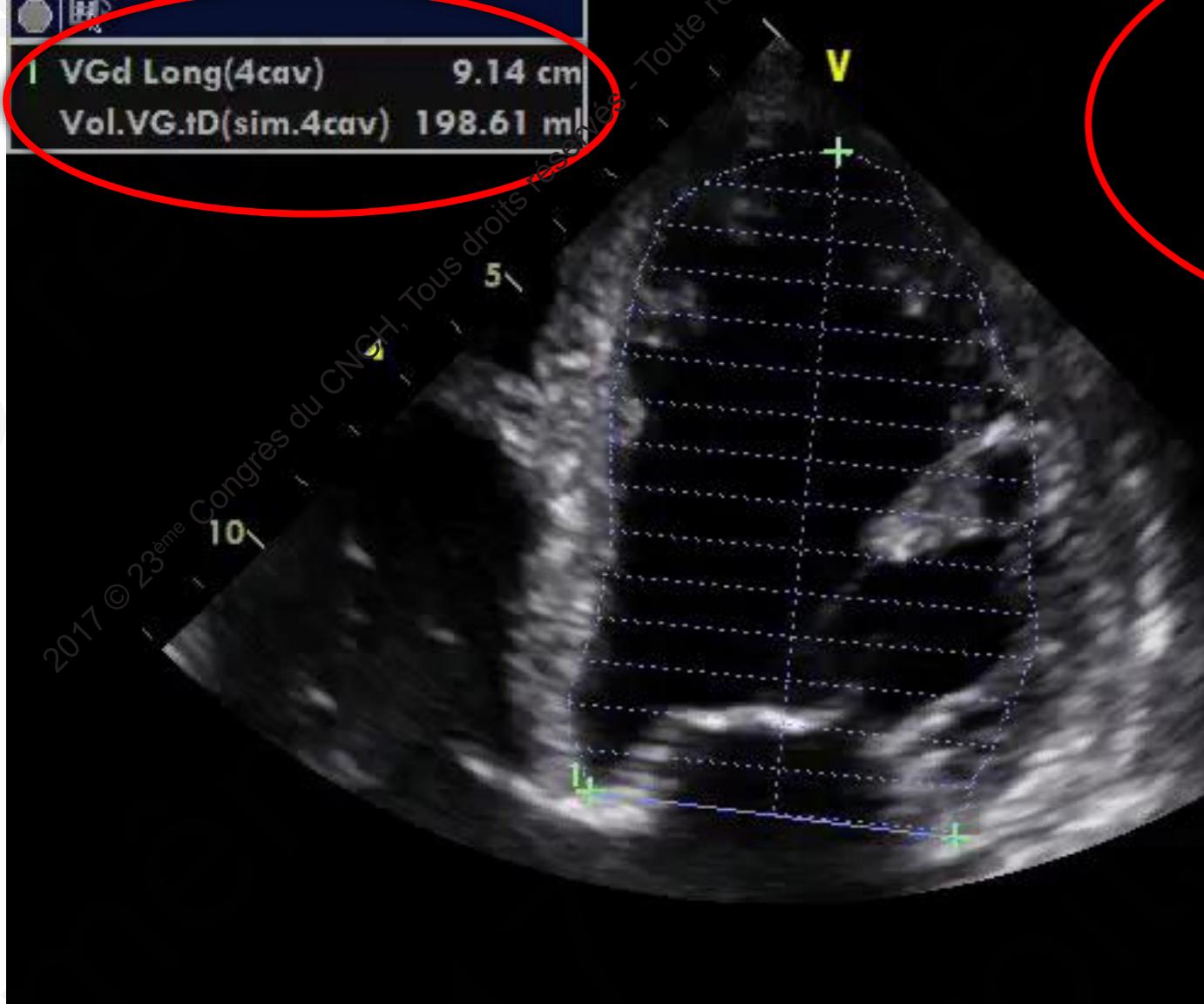
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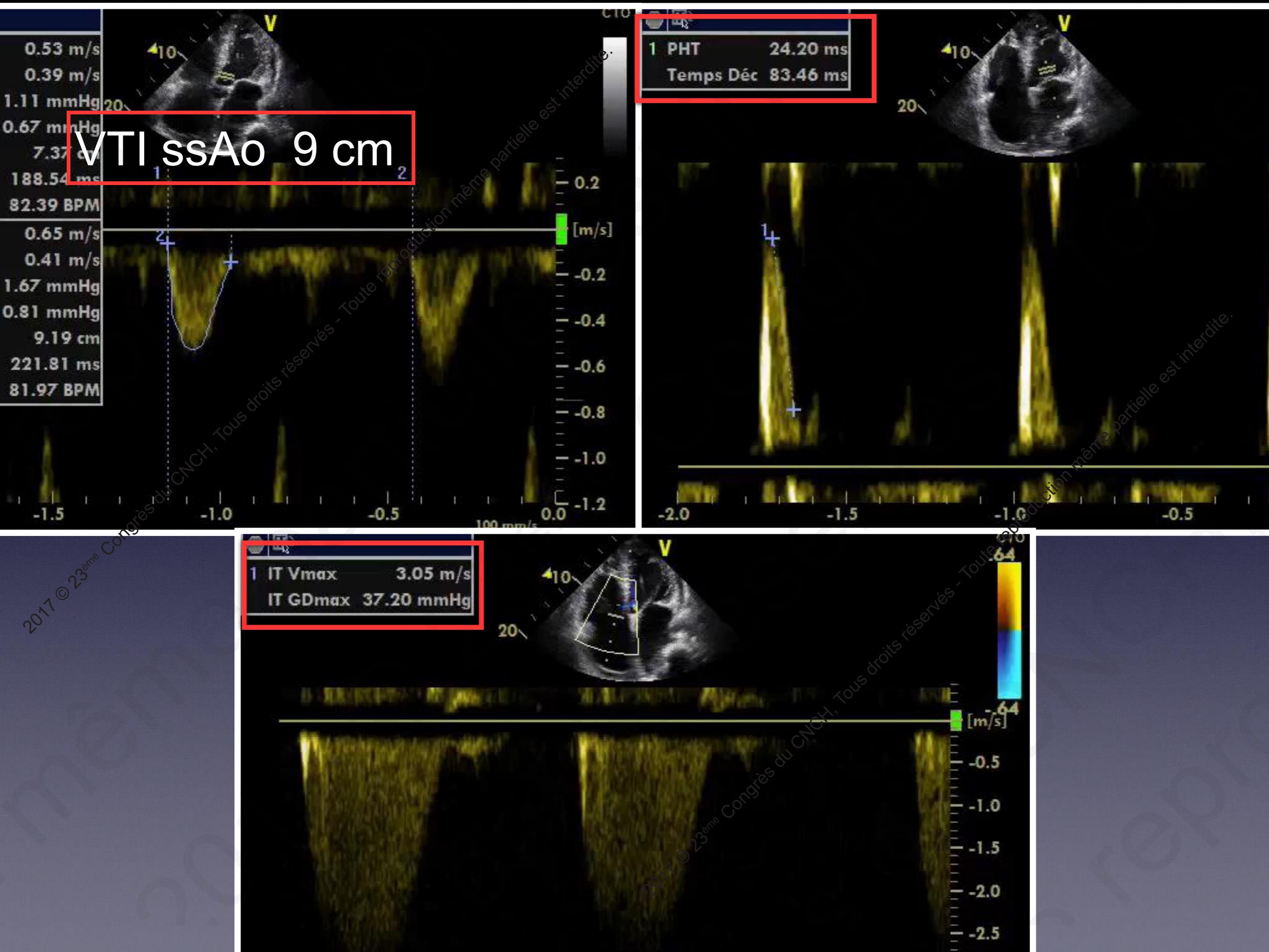
ESC 2016

DYSPNEE aigue



DYSPNEE aigue





DYSPNEE aigue

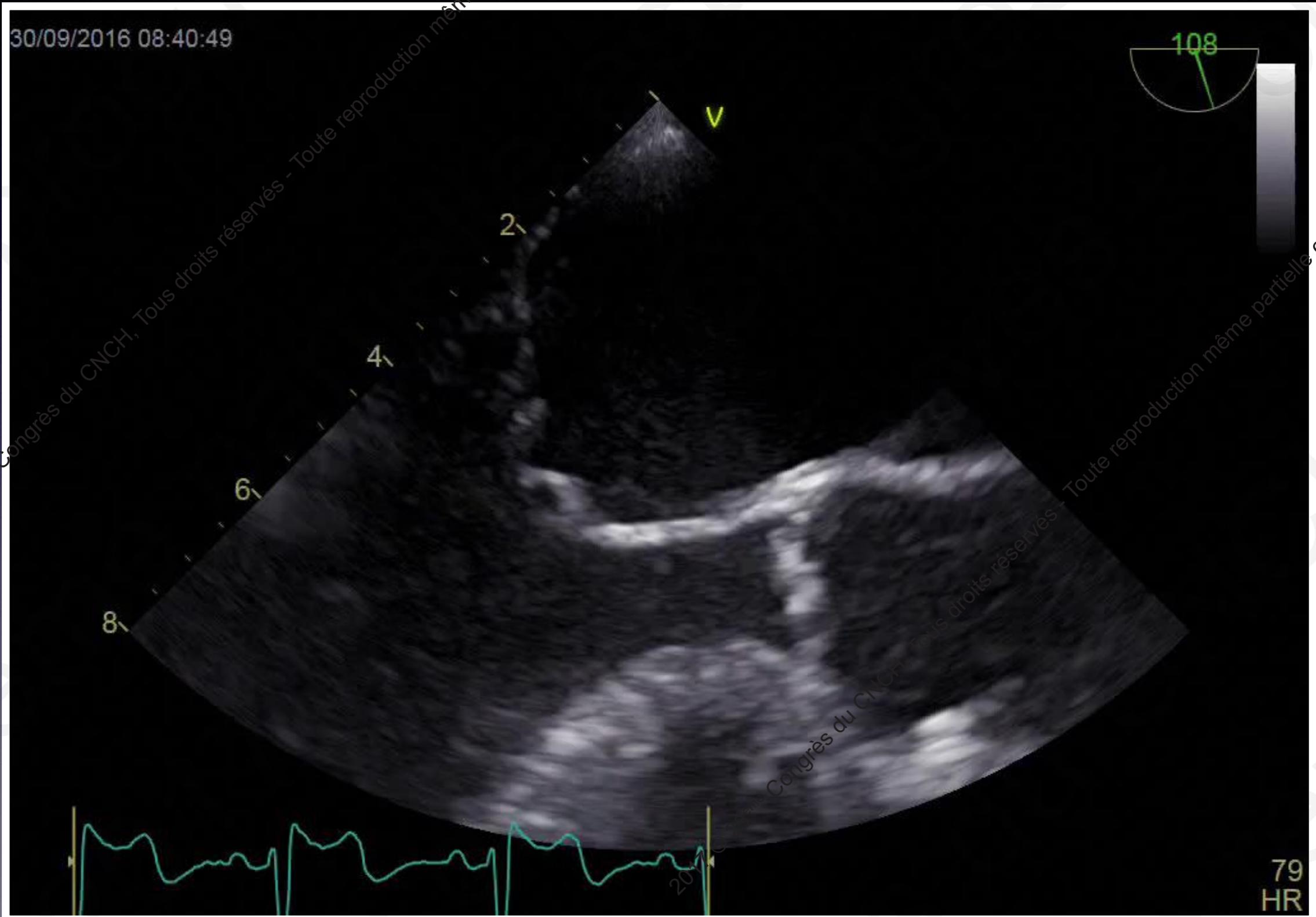
Dysfonction systolique majeure avec baisse du débit cardiaque:

Diagnostic facile

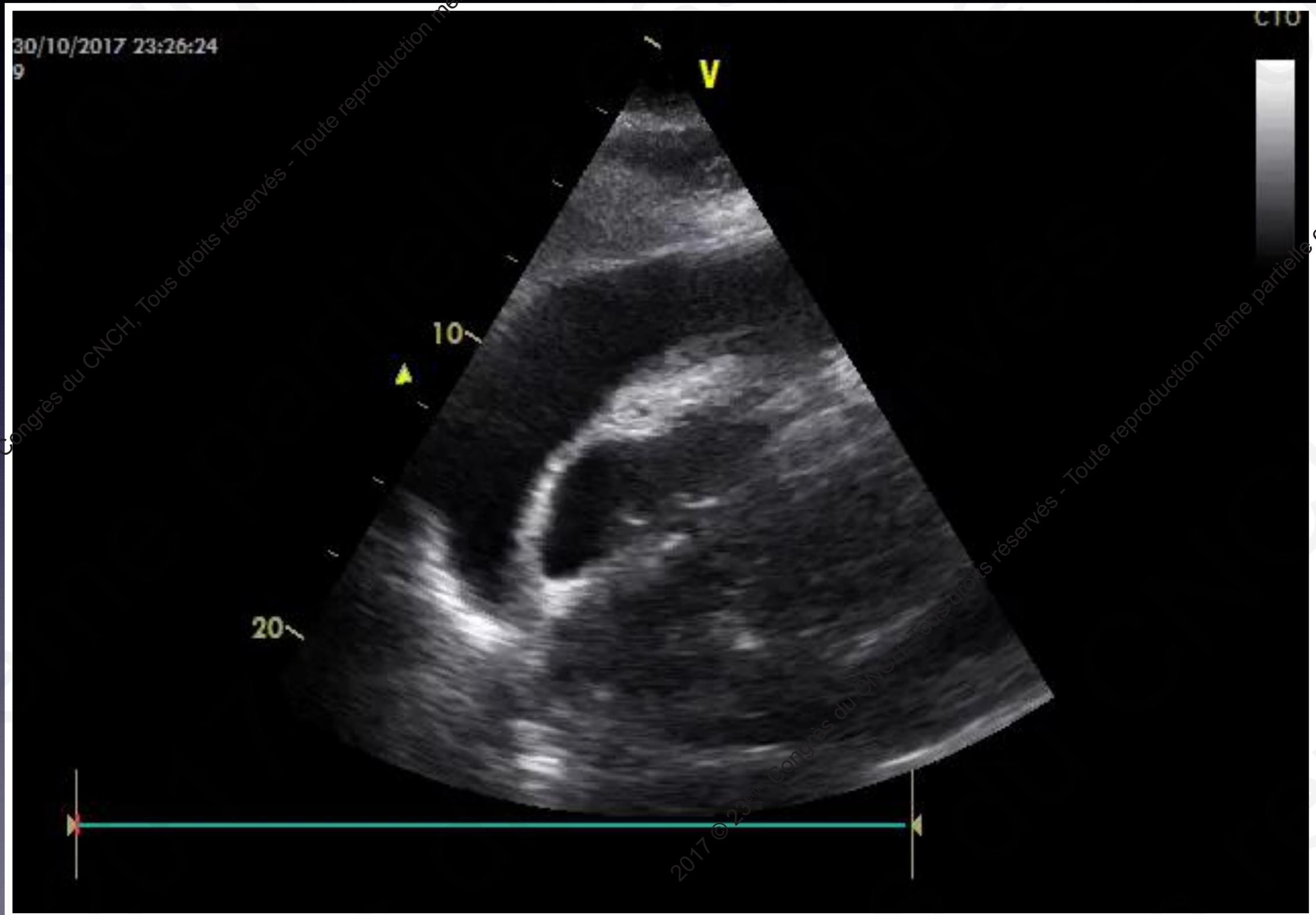
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L'ECHO c'est facile



L'ECHO c'est facile

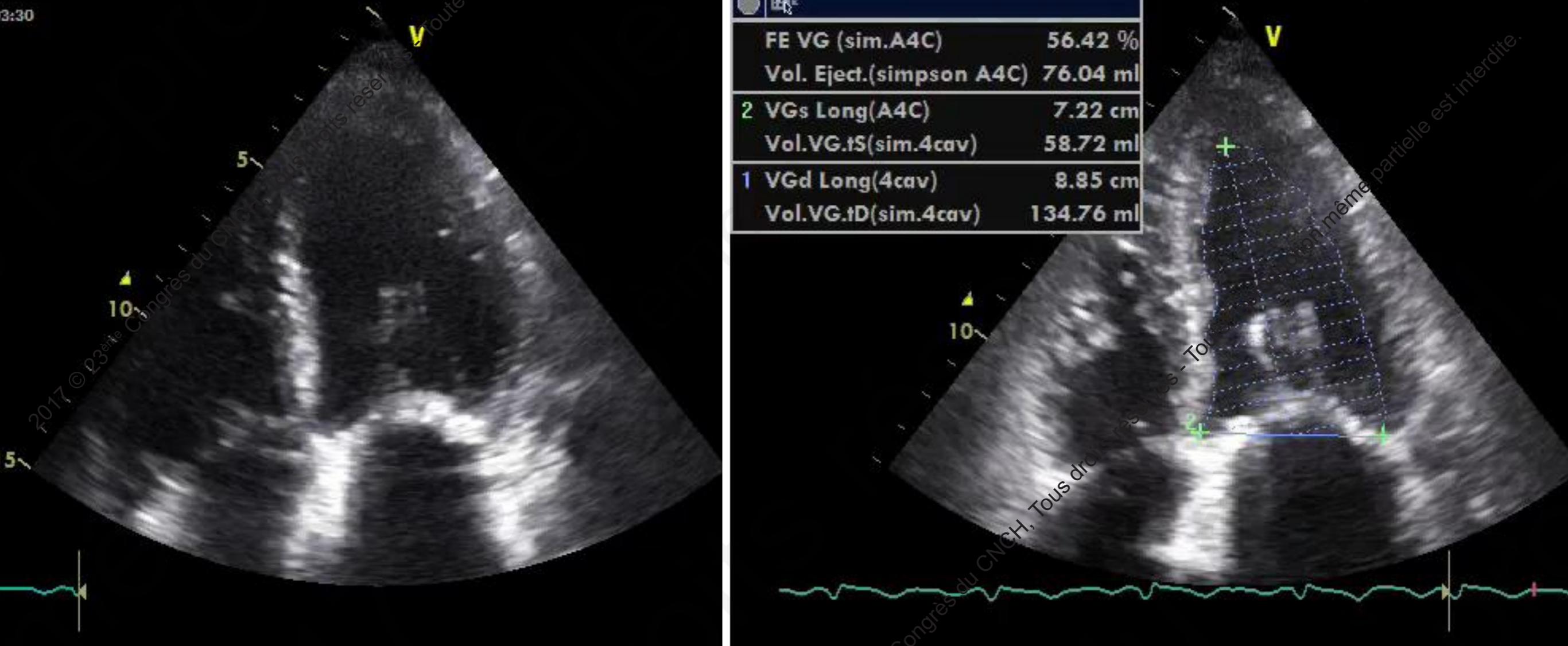


Mais pas toujours.

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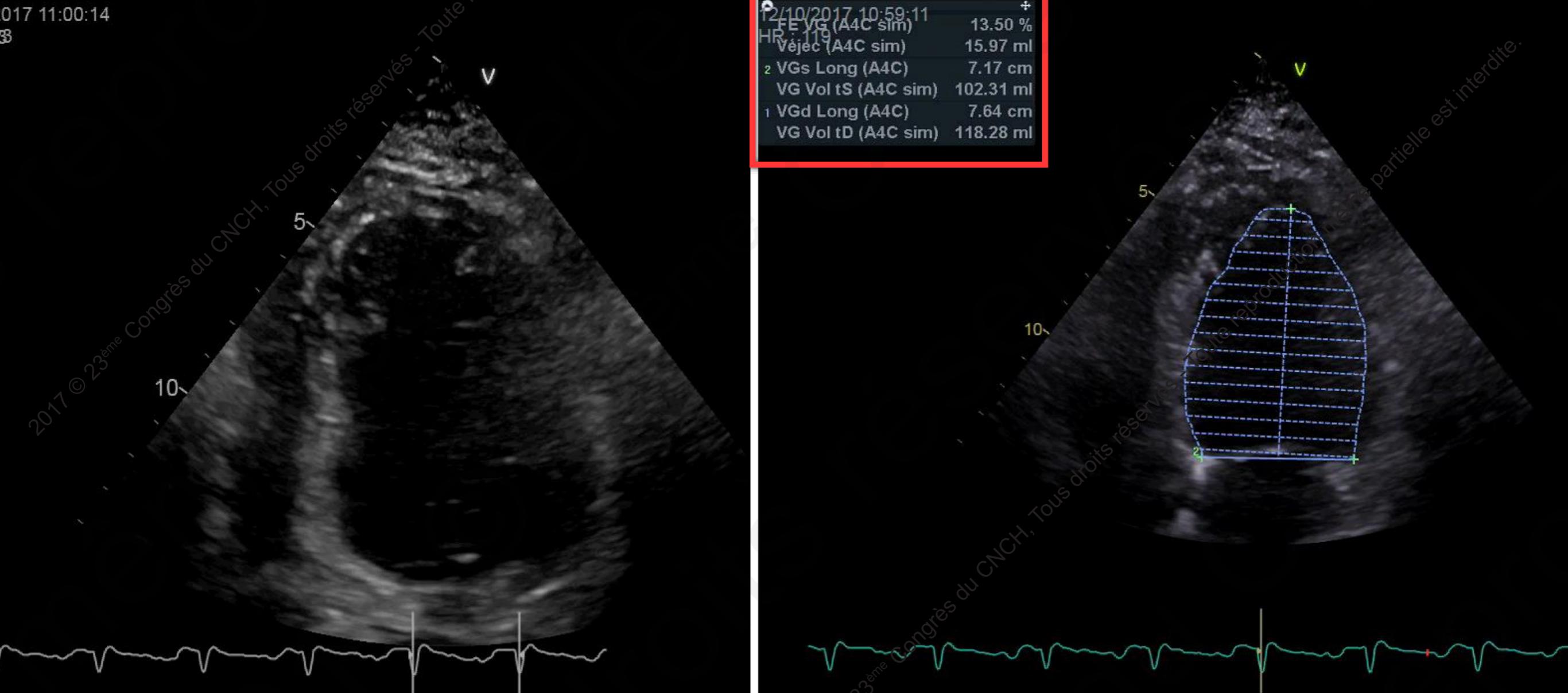
Patient n°1



Patient n°2

2017 11:00:14

3



**LEQUEL DE CES PATIENTS EST ADMIS
EN USIC et MALHEUREUSEMENT N'EN
SORTIRA PAS,
PENDANT QUE L'AUTRE GAMBADE DANS
LA NATURE ?**

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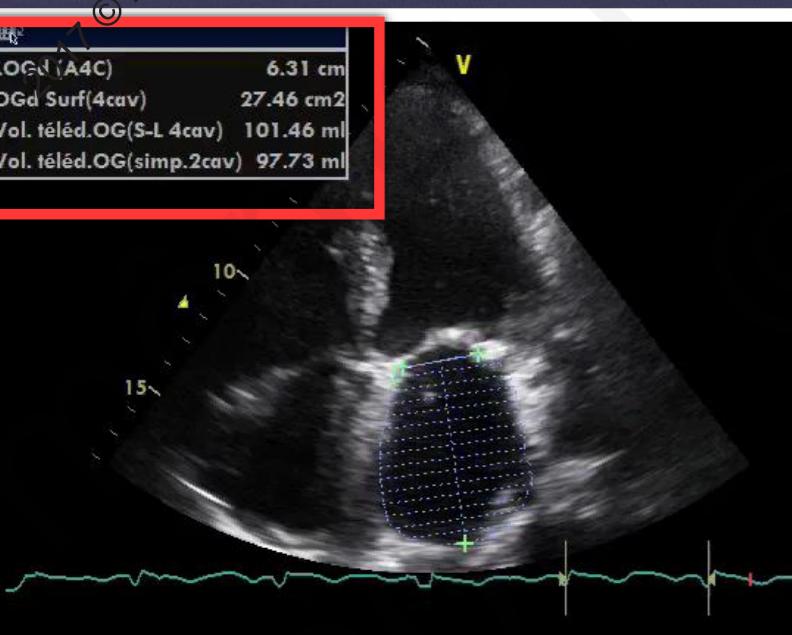
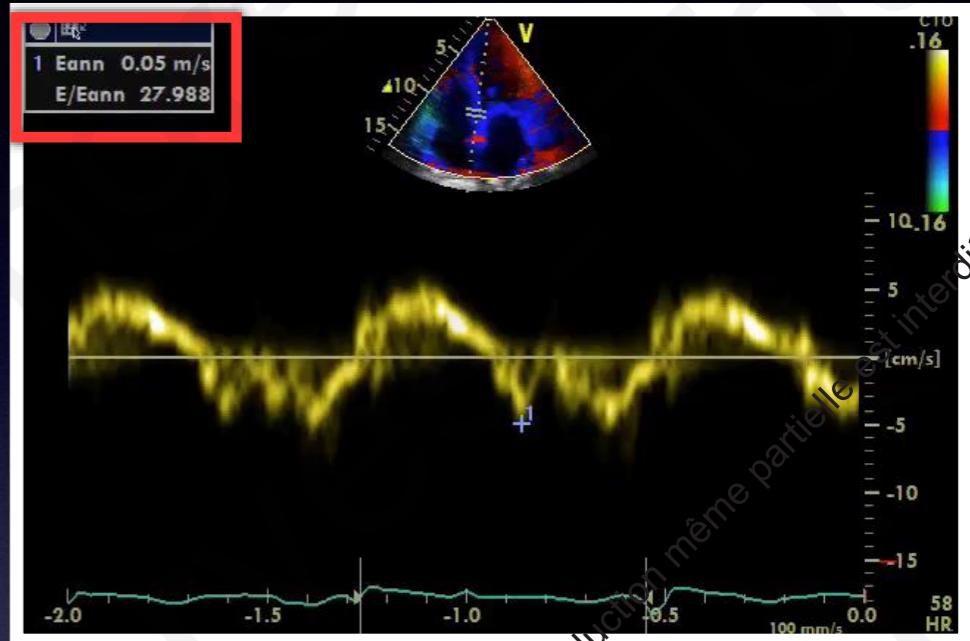
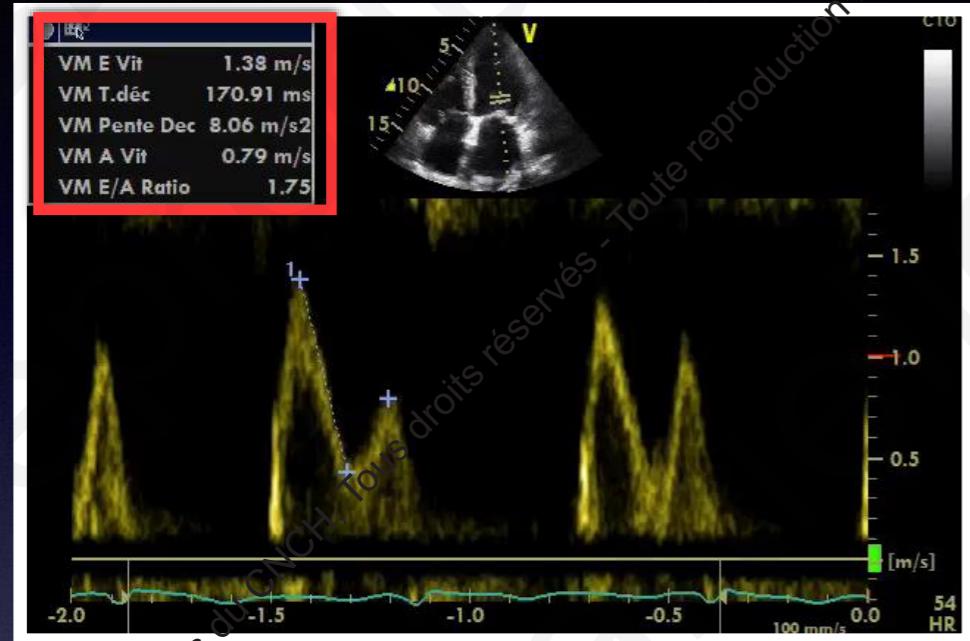
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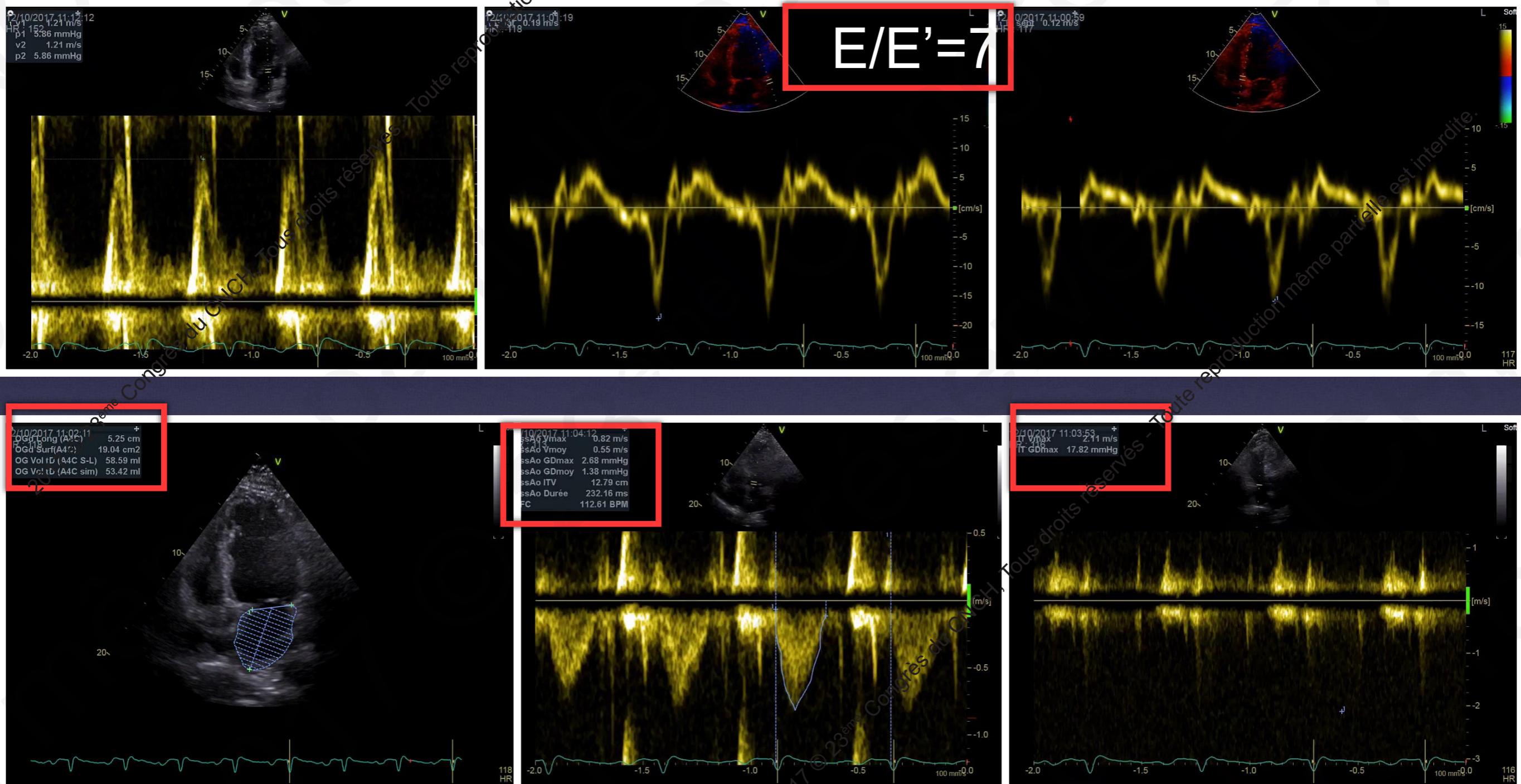
Patiente n°1

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Patient n°1



Patient n°2



ICA à FE préservée

- Def: Tableau d'ICA avec une FE > 40-50%
- Au moins 45% des ICA
- Ratio femme/homme 3/2
- HTA ou Card. Isch ou Diabète > 90%
- Histoire d'ICA : > 60%
- Pronostic (presque) aussi mauvais que FE altérée

ICA à FE préservée

Prévalence

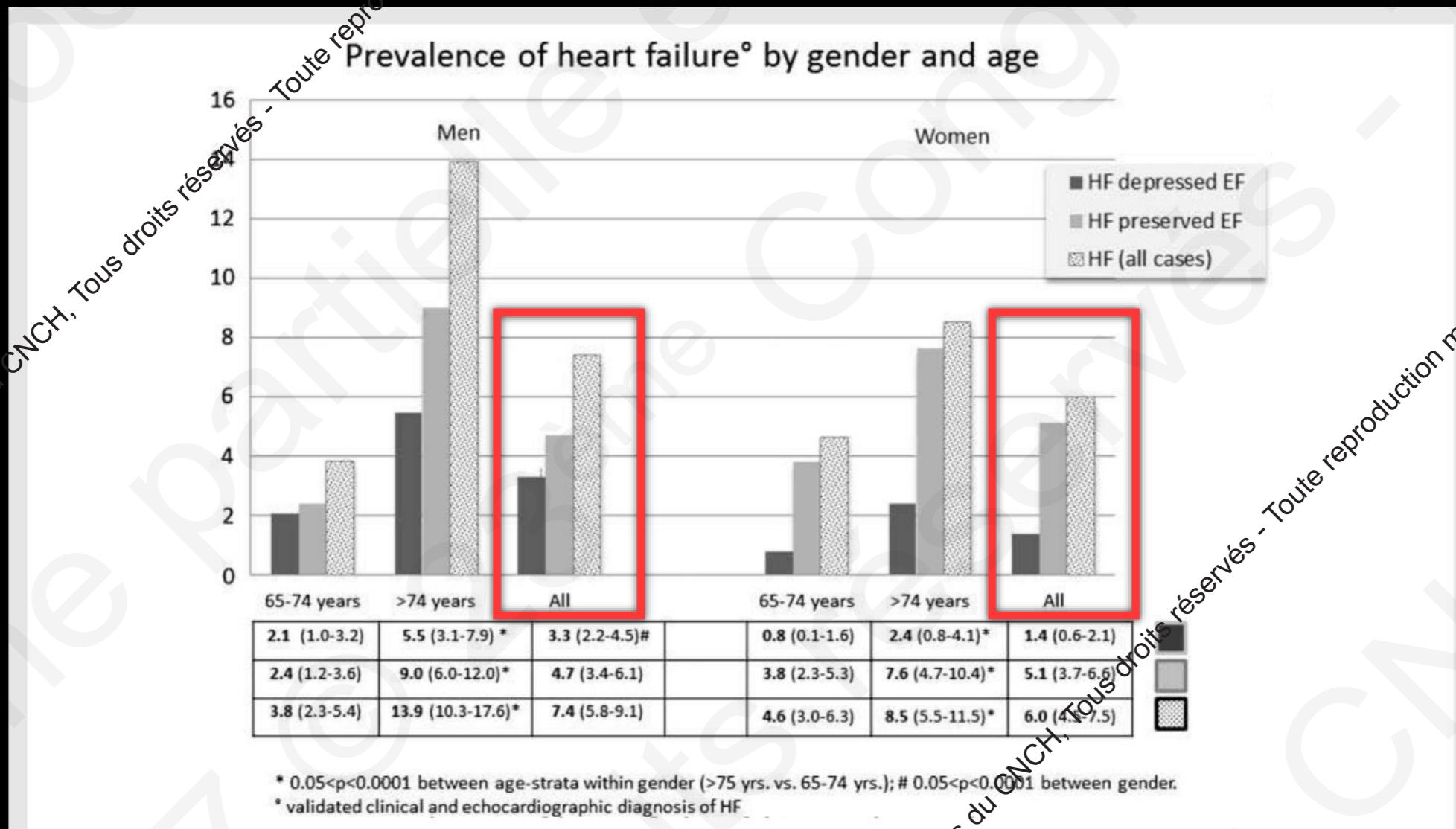


Figure 2 Prevalence of heart failure (HF) by gender and age. Prevalence of HF as a percentage (ordinate, %) is shown for age (65–74 years, >75 years, and all) and gender groups (abscissa). Dark grey bars, HF with depressed ejection fraction (EF); light grey bars, HF with preserved EF; dotted bars, all cases of HF. Mean percentage (in bold) and the 95% confidence interval (in parentheses) are shown for each condition (HF depressed EF, HF preserved EF, and all cases) at the bottom of the figure^o

IC à FE préservée Sous diagnostiquée

Table 2 Prevalence of previously unknown heart failure with reduced ejection fraction (HF-REF) and heart failure with preserved ejection fraction (HF-PEF) stratified by age and sex

Age (years)	Male		Female		Total	
	n	HF-REF	n	HF-REF	n	HF-REF
64–74	155	5 (3.2)	179	2 (1.1)	334	7 (2.1)
75–84	111	6 (5.4)	140	4 (2.9)	251	10 (4.0)
All ages	266	11 (4.1)	319	6 (1.9)	585	17 (2.9)
95% CI	2.2–7.5	7.9–15.9	0.8–4.3	9.2–16.8	1.8–4.7	9.5–14.9
						All HF*
						28 (8.4)
						64 (25.5)
						92 (15.7)
						12.9–19.0

*In all heart failures; the five cases of isolated right-sided heart failure were also included.

Values are numbers (percentage); 95% CI, 95% confidence interval.

ICA à FE préservée Prévalence

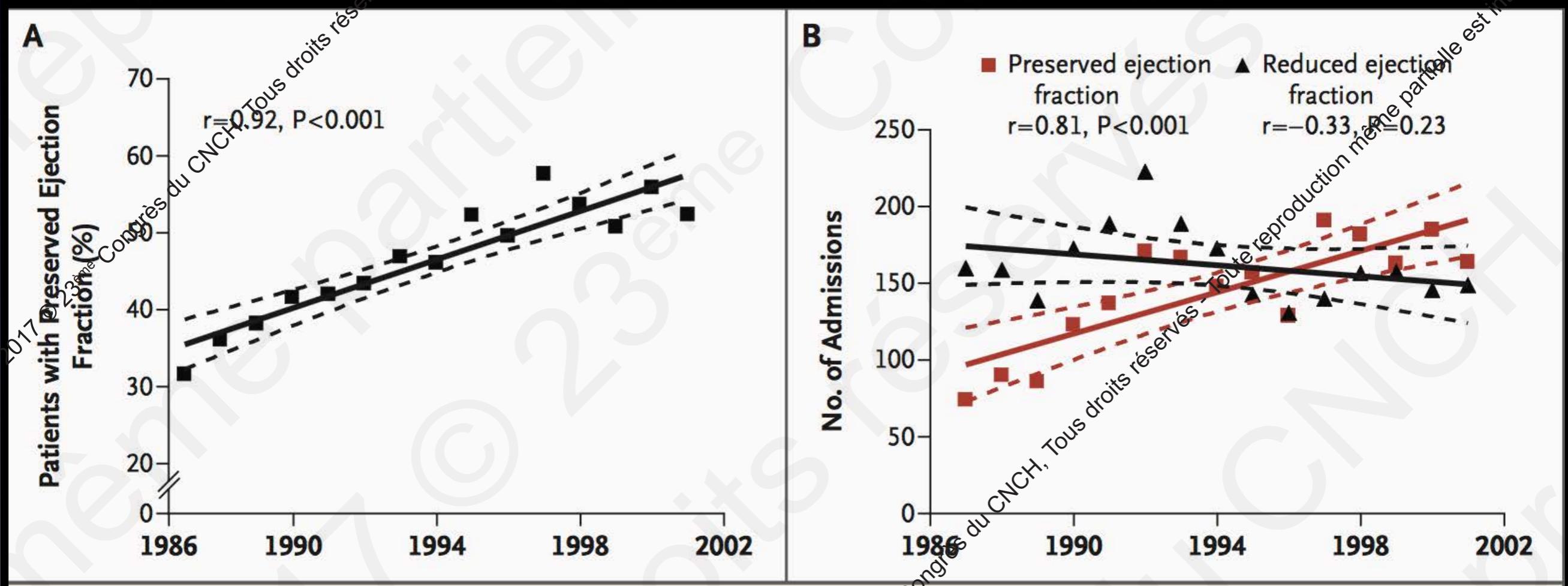
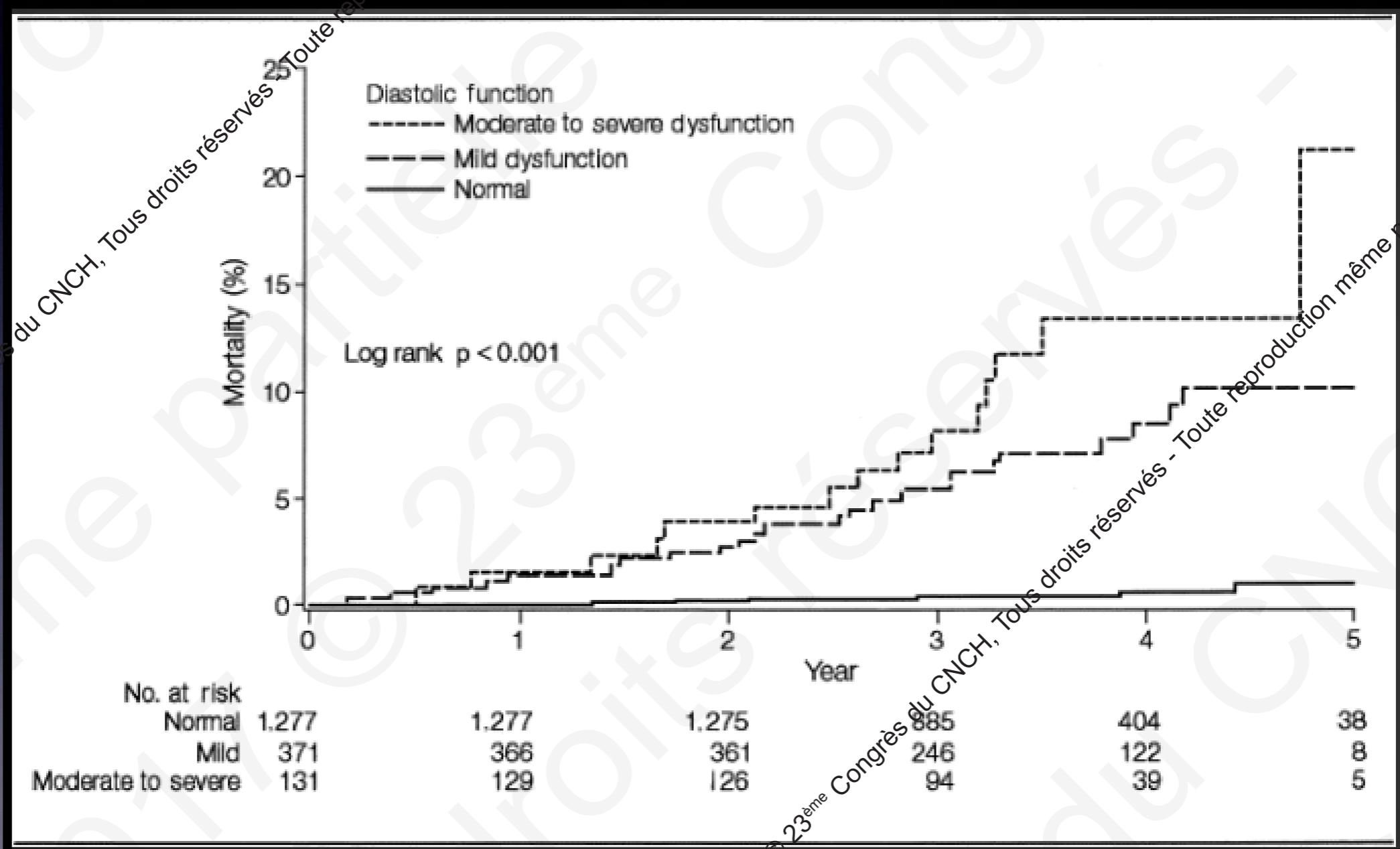


Figure 1. Secular Trends in the Prevalence of Heart Failure with Preserved Ejection Fraction.

IC à FE préservée Pronostic



IC à FE préservée Pronostic

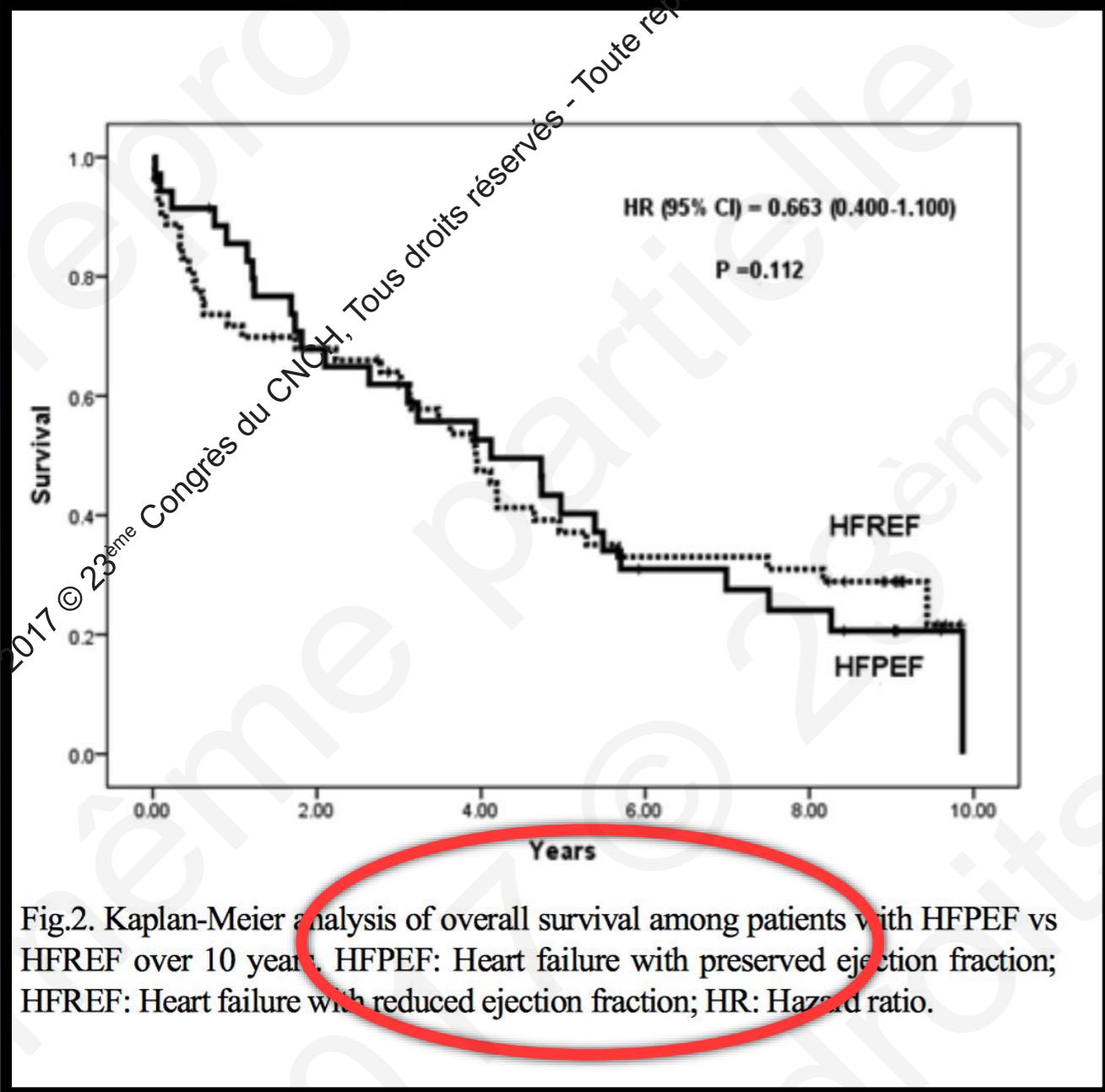


Fig.2. Kaplan-Meier analysis of overall survival among patients with HFPEF vs HFREF over 10 years. HFPEF: Heart failure with preserved ejection fraction; HFREF: Heart failure with reduced ejection fraction; HR: Hazard ratio.

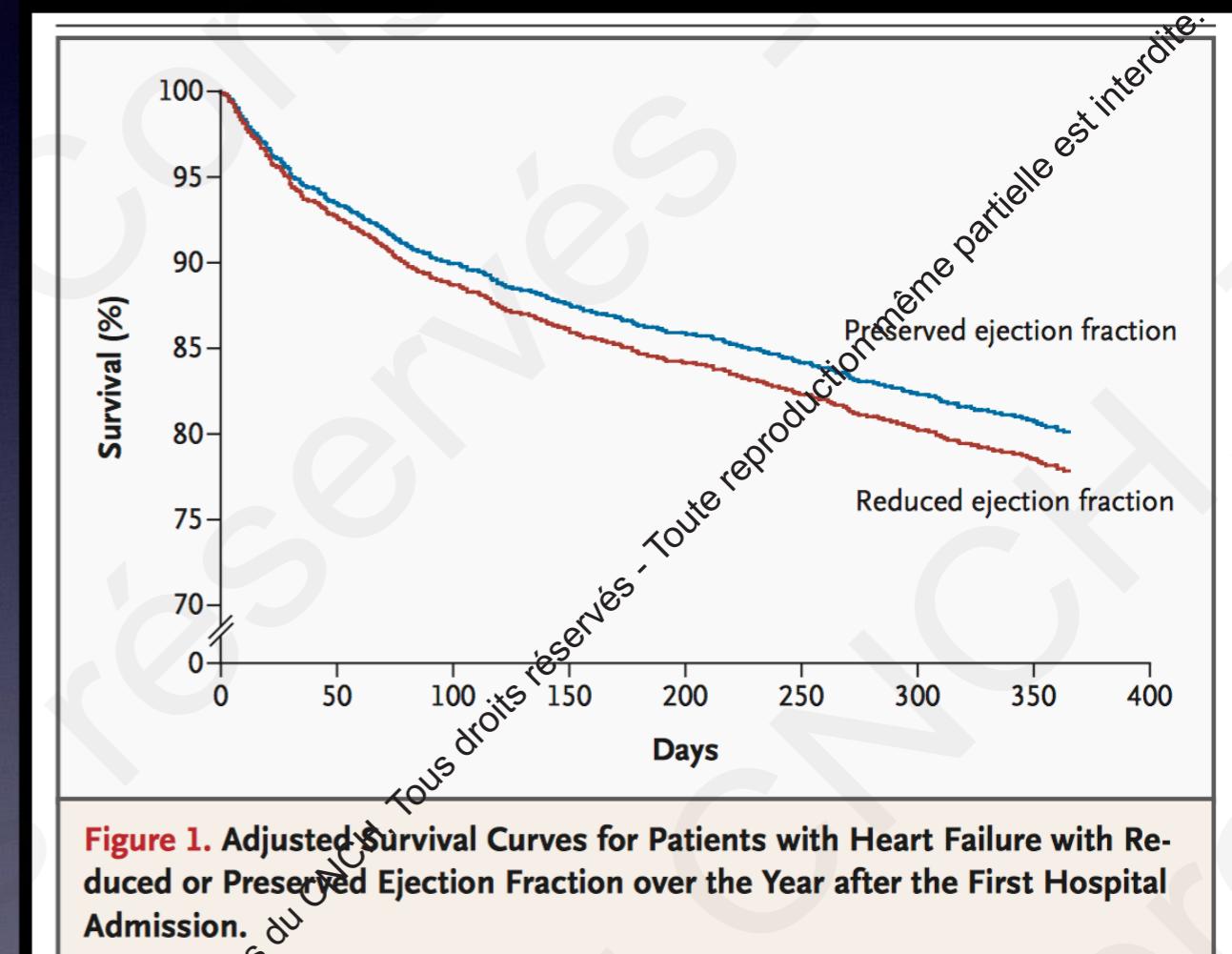


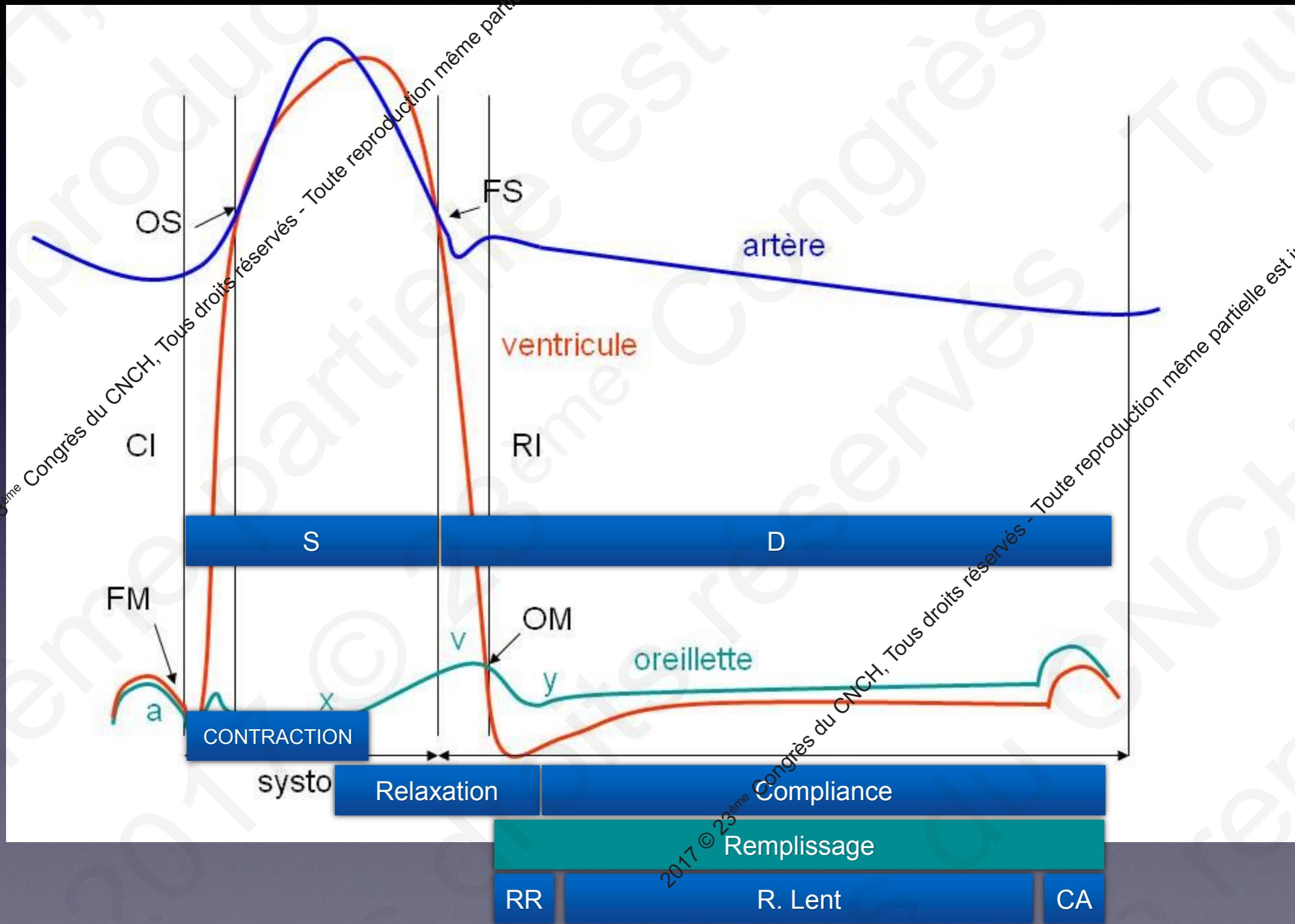
Figure 1. Adjusted Survival Curves for Patients with Heart Failure with Reduced or Preserved Ejection Fraction over the Year after the First Hospital Admission.

COMMENT (BIEN) EVALUER CES PATIENTS ?

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Diastologie en bref

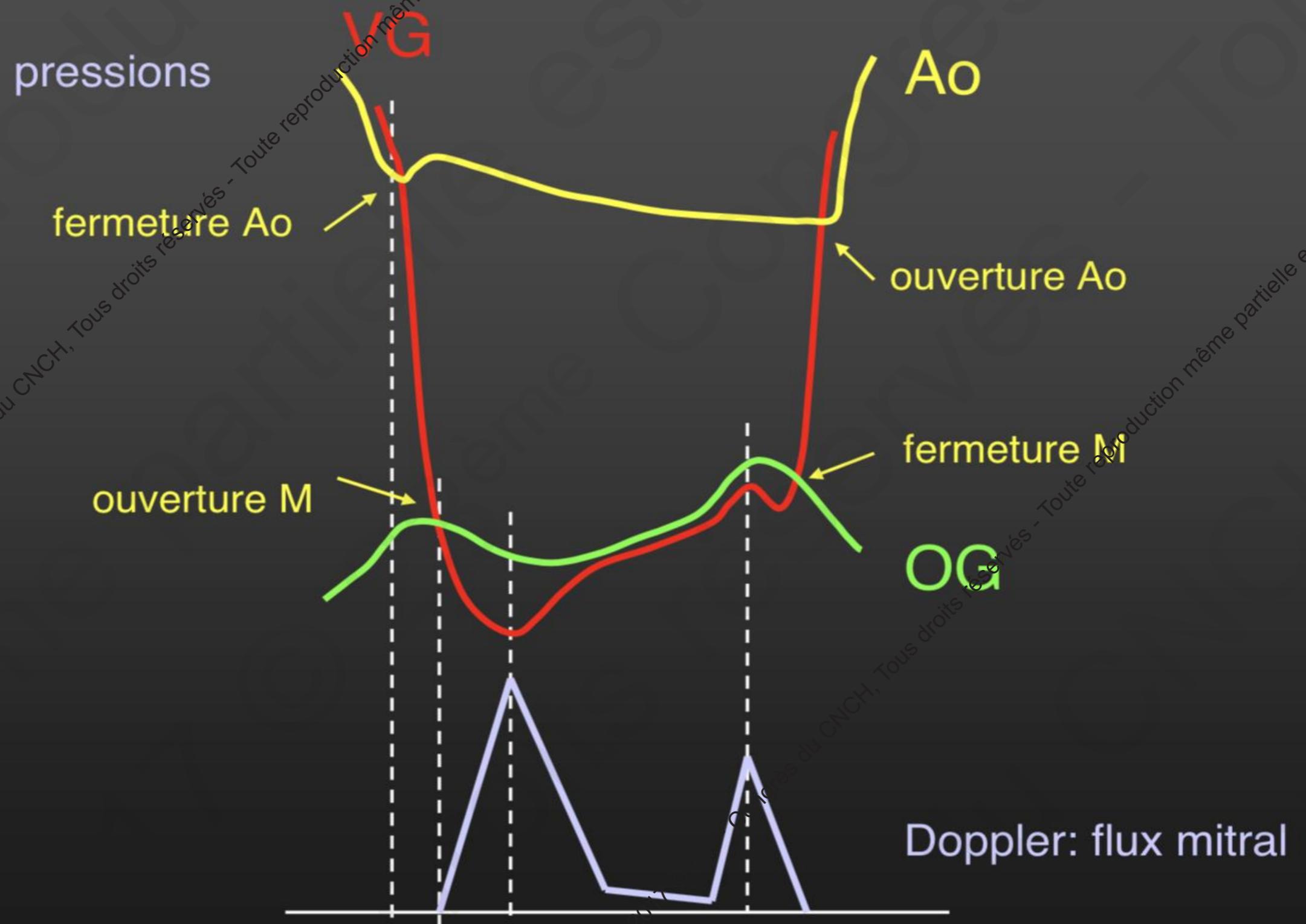


Paramètres à recueillir

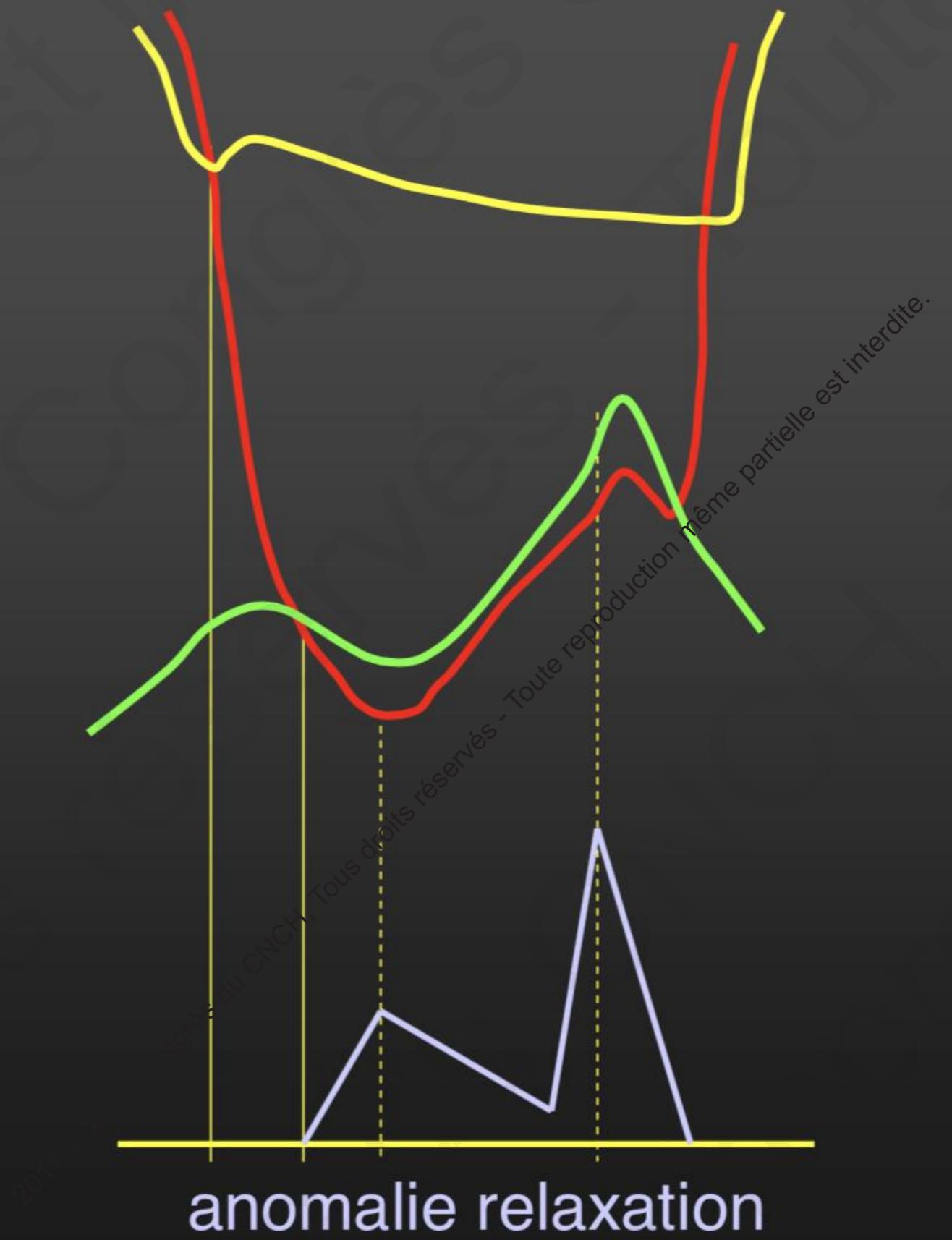
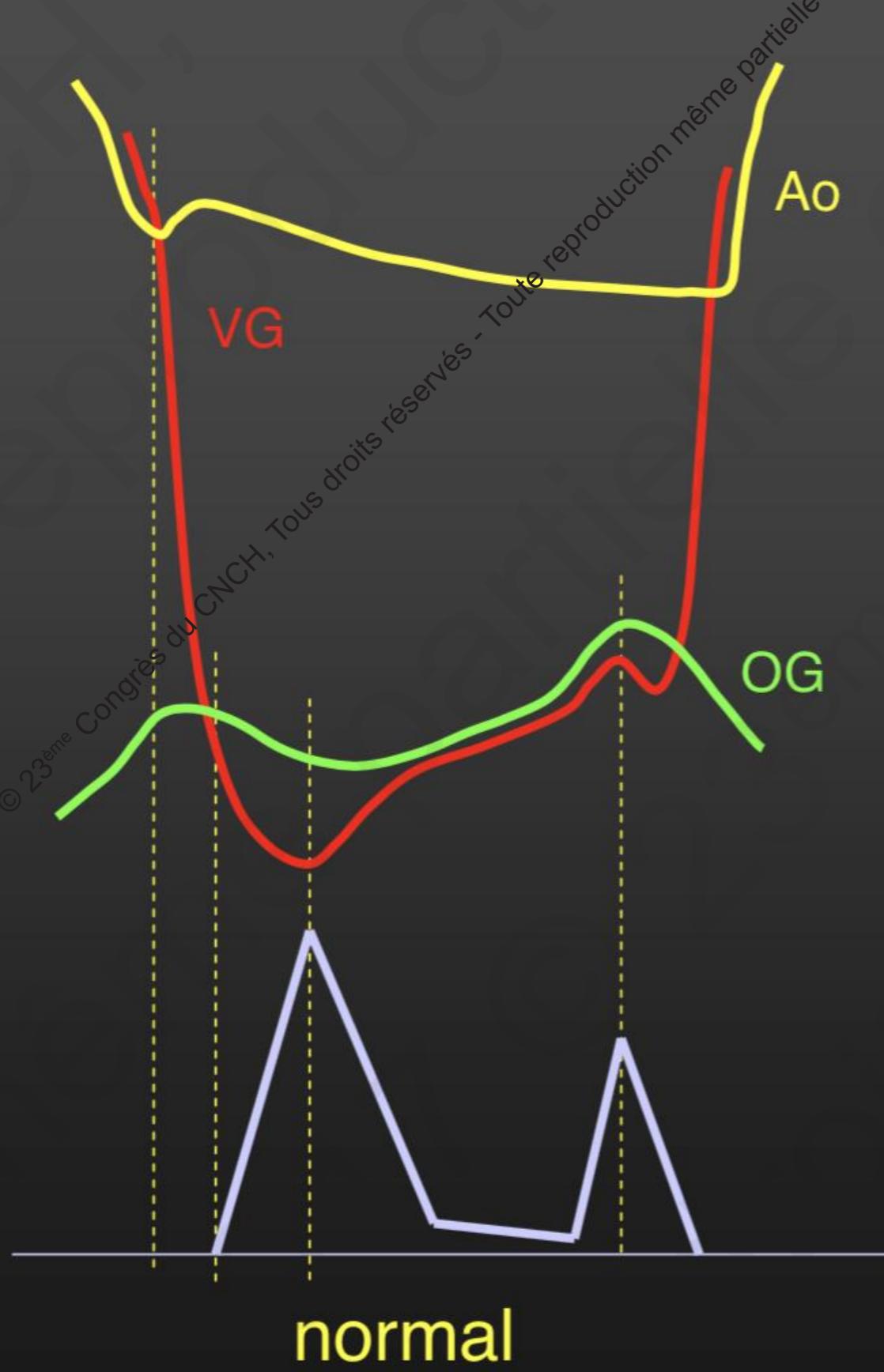
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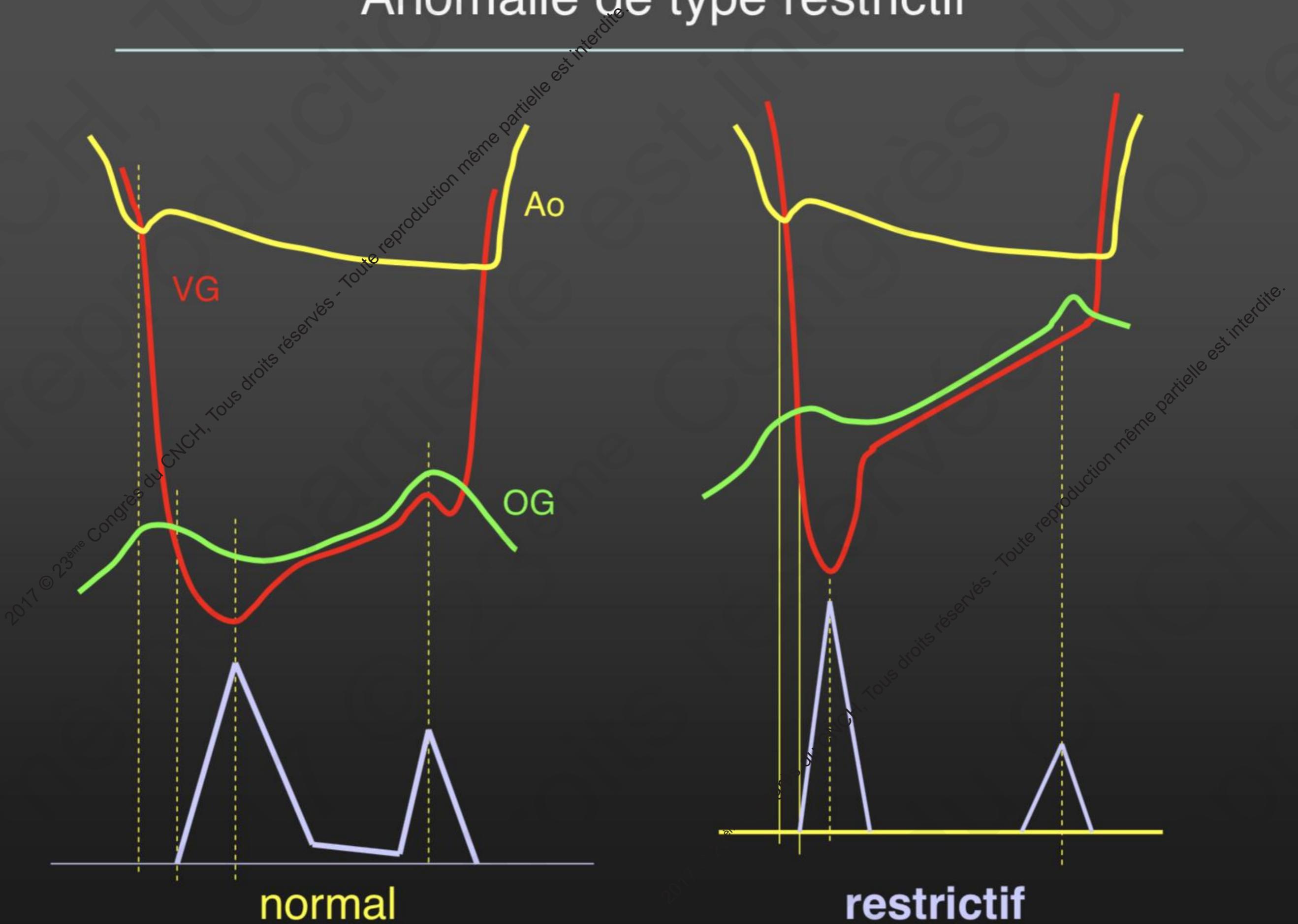
Flux transmitral normal



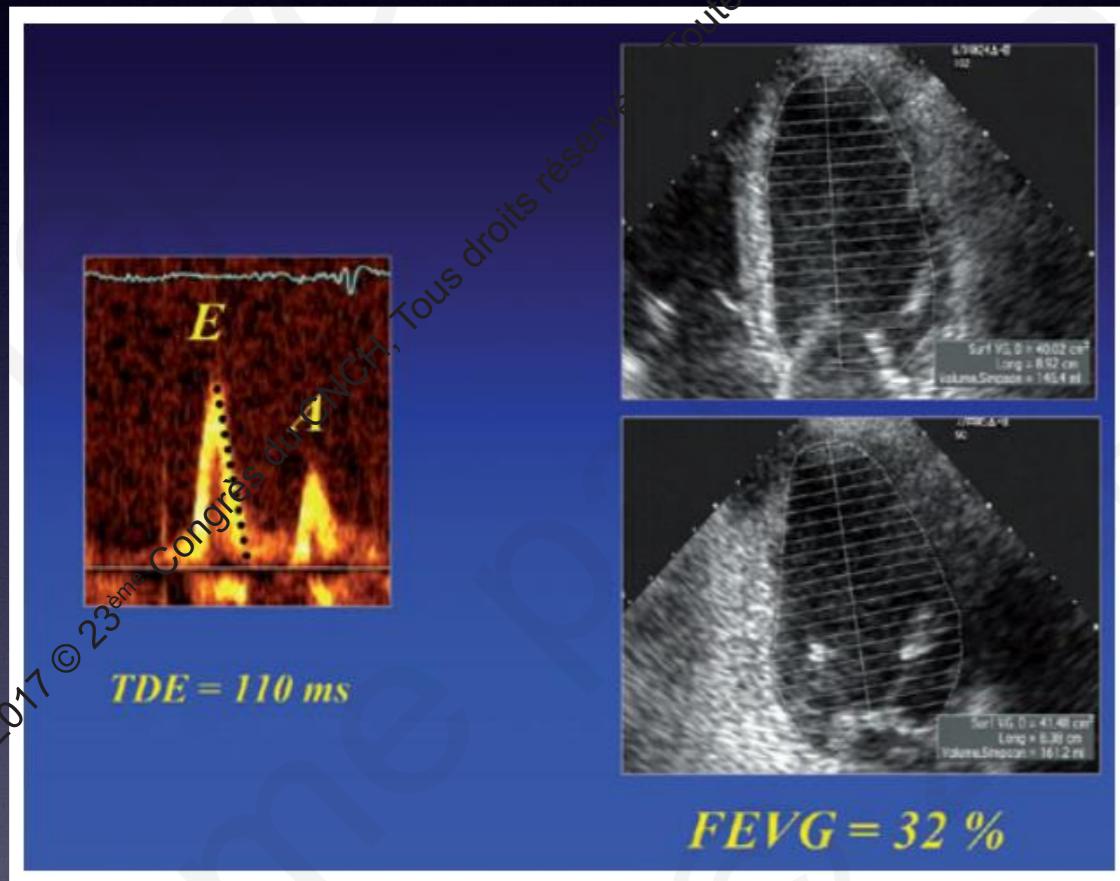
Anomalie de relaxation



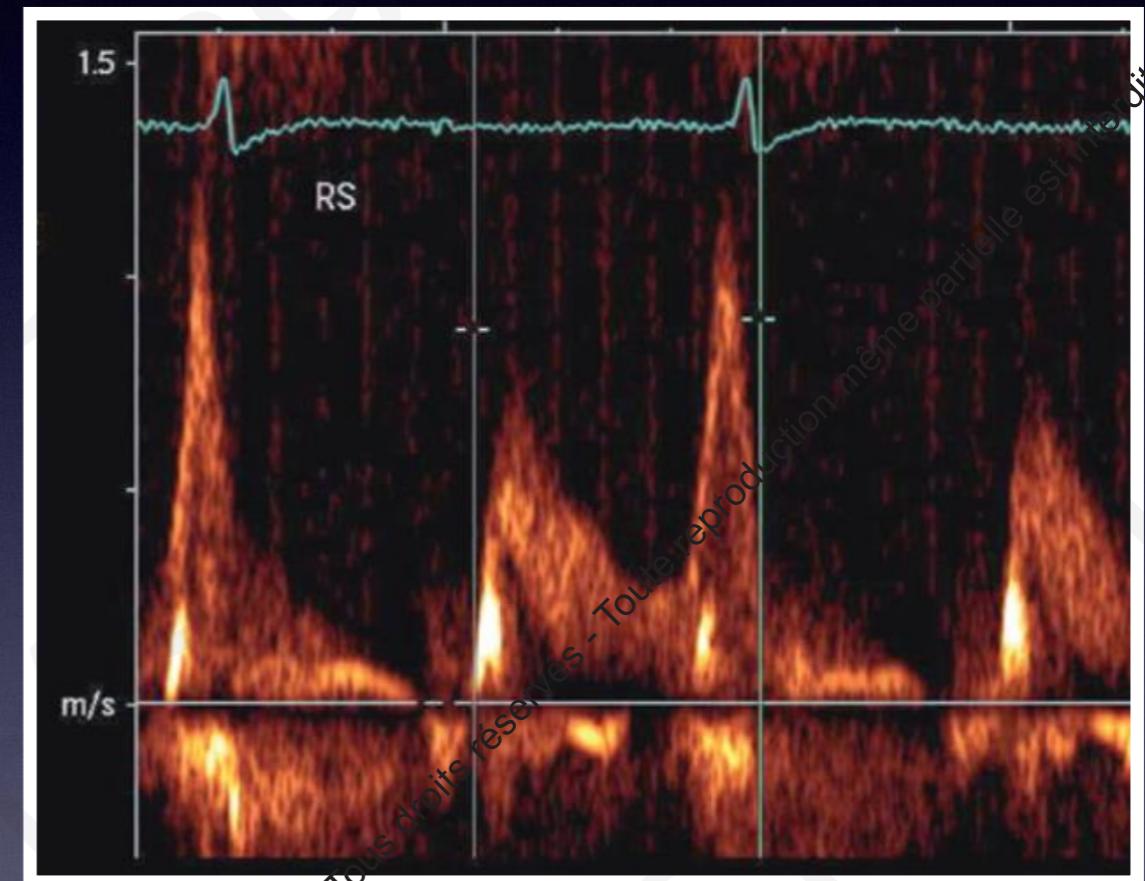
Anomalie de type restrictif



Peut on se contenter du flux mitral ?

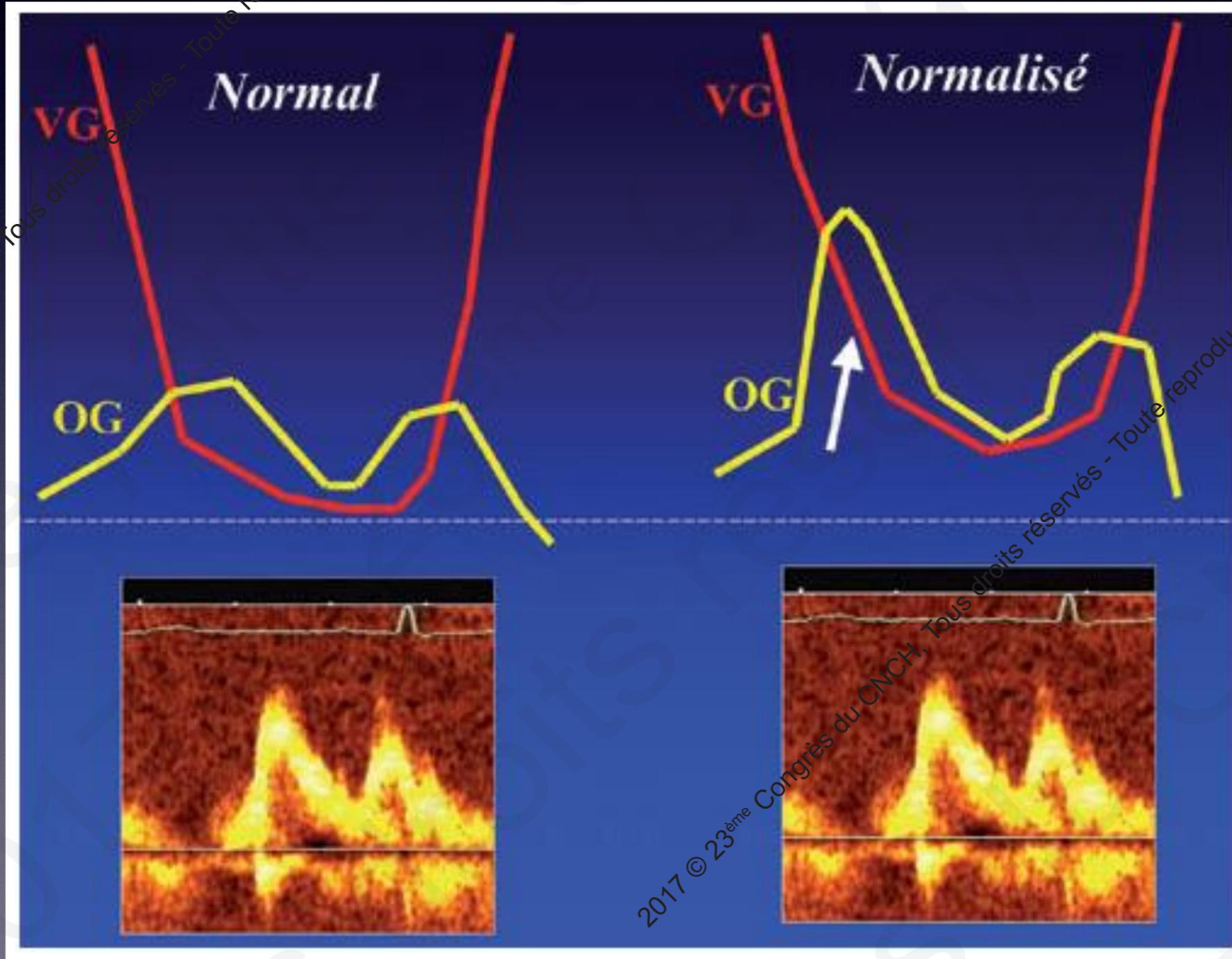


PRVG toujours élevées
(sauf grand sportif)

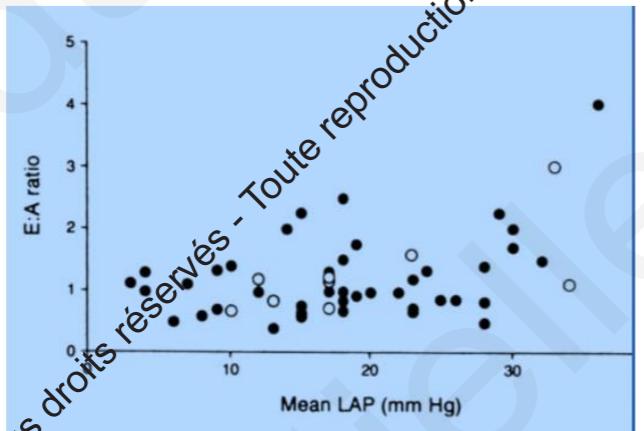


PRVG presque tjs basses
(sauf si CMH)

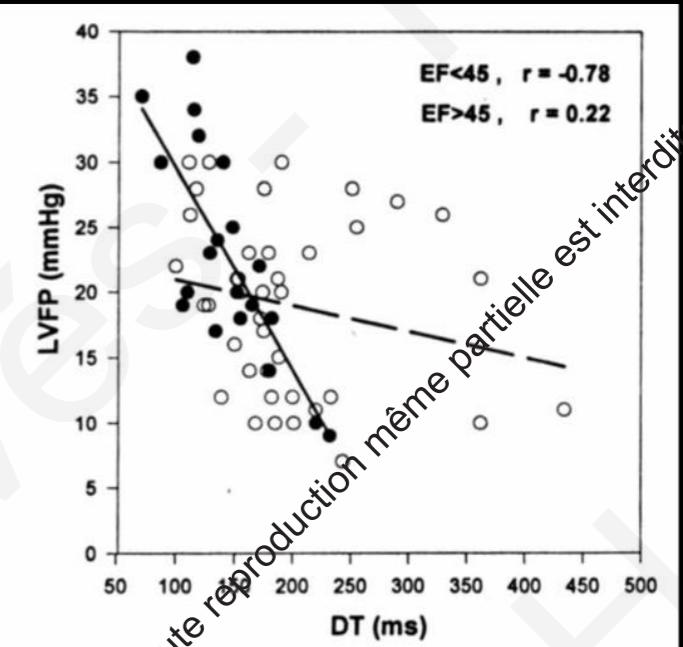
Le plus souvent



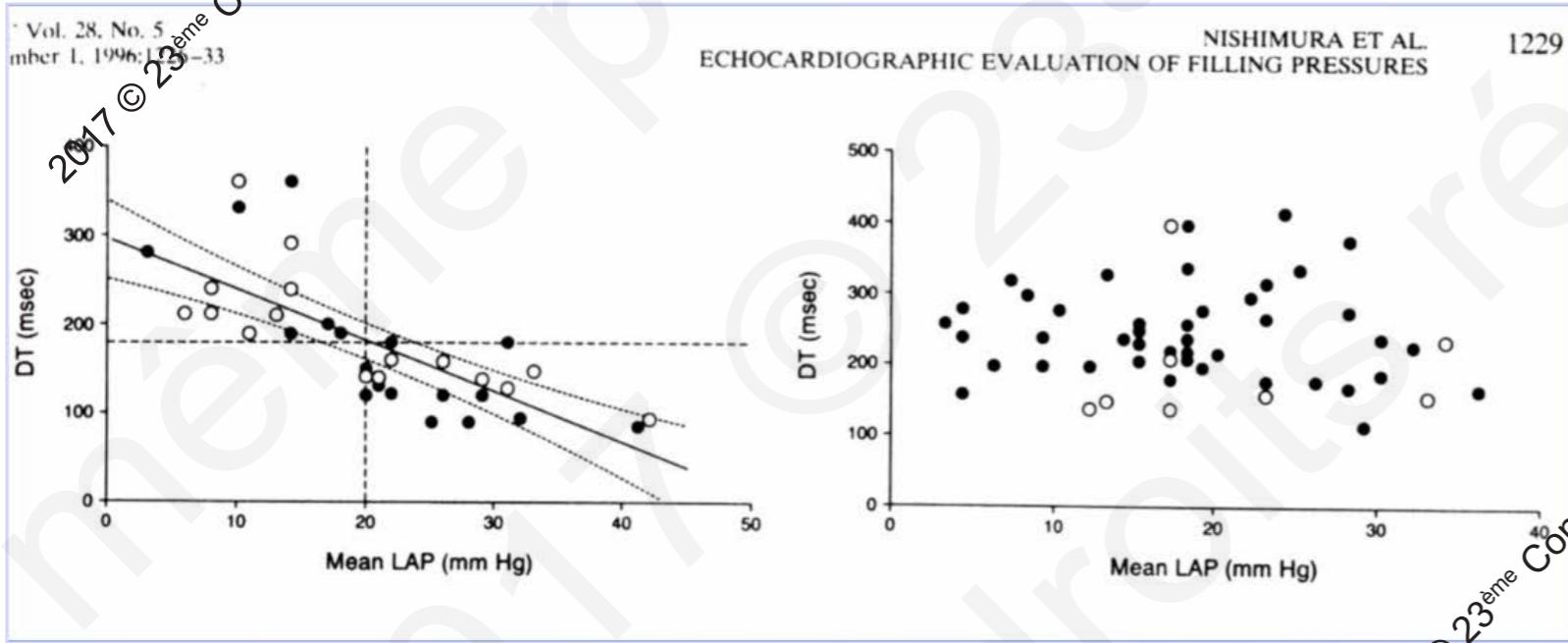
Peut on se contenter du flux mitral ?



Mauvaise corrélation entre POG et ratio E/A
(Nishimura et al JACC 1996; 5: 1226-33)

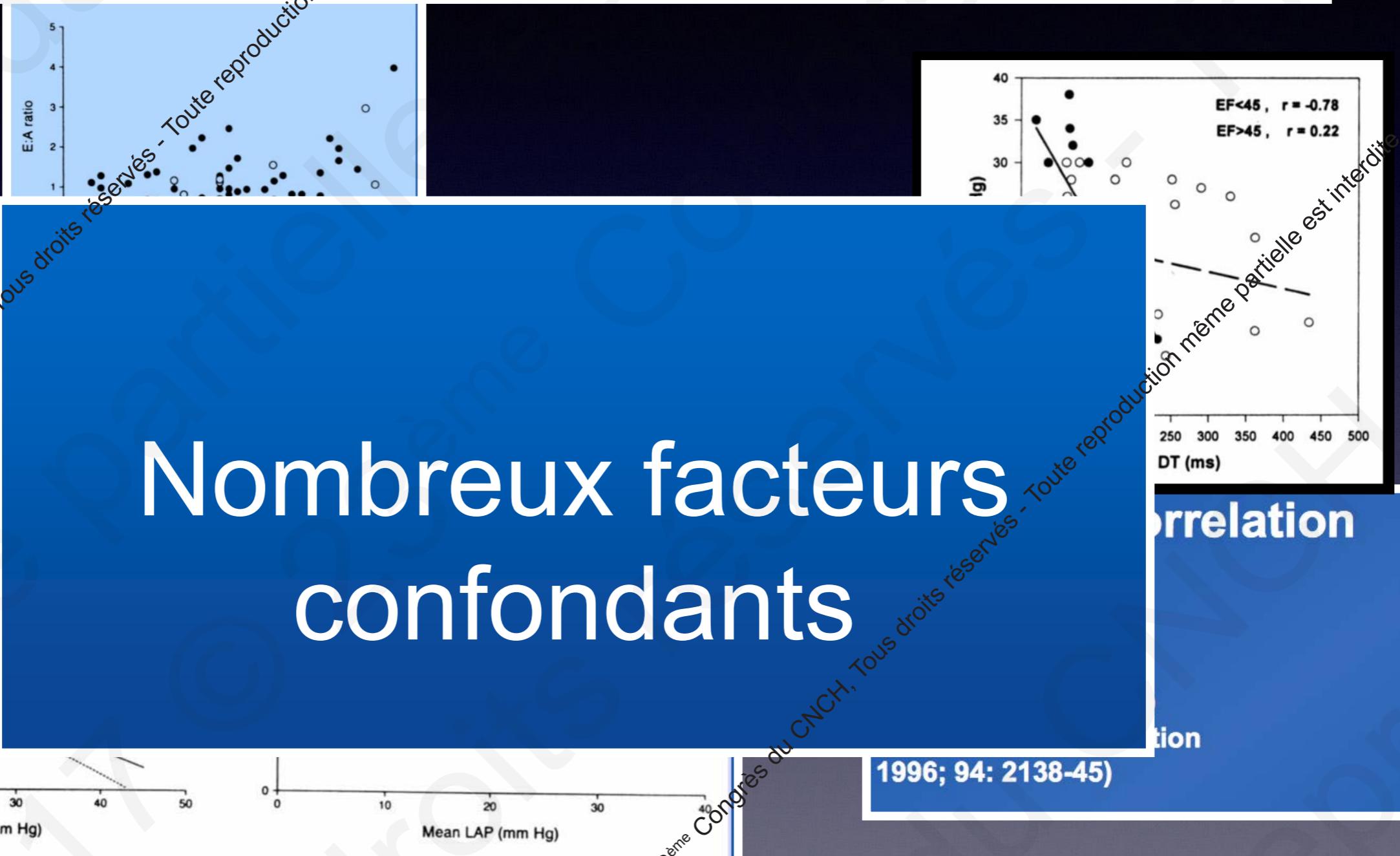


Mauvaise corrélation entre TDE PTDVG si FE > 45 %
(Nagueh et al Circulation 1996; 94: 2138-45)



Mauvaise corrélation entre POG et TDE
(Nishimura et al JACC 1996; 5: 1226-33)

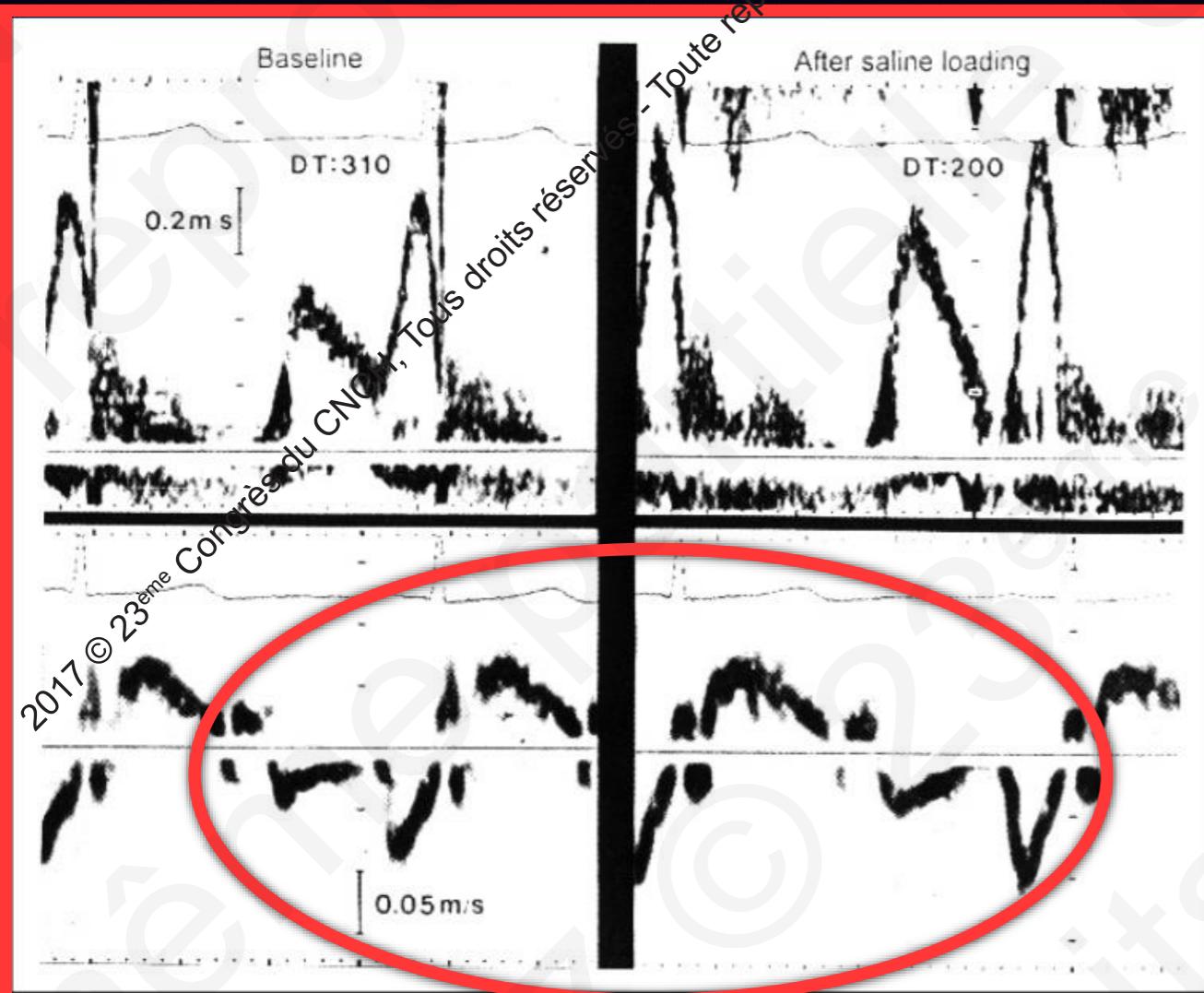
Peut on se contenter du flux mitral ?



Facteurs de variations du flux mitral

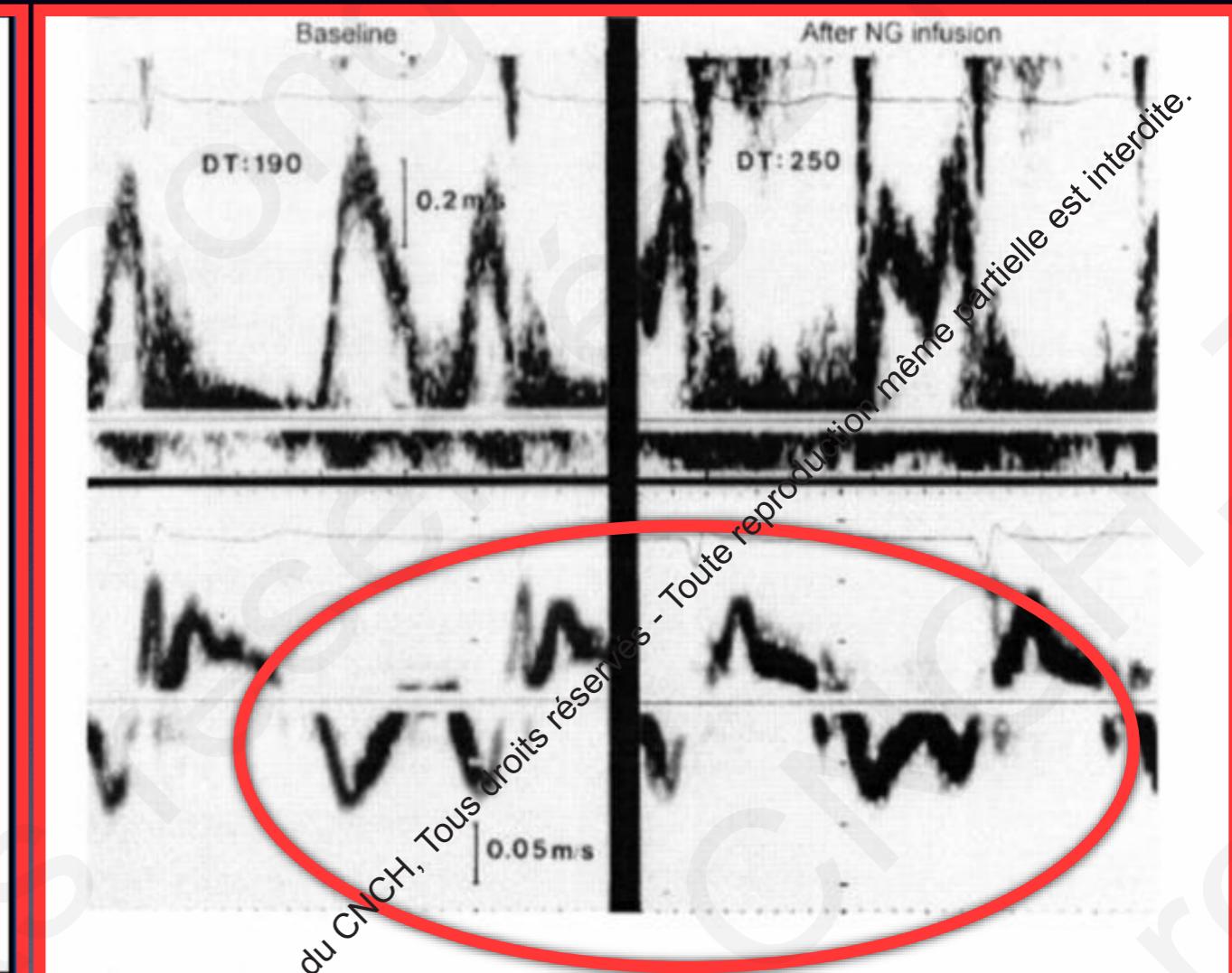
- Age
- Fréquence cardiaque
- Conduction AV et InterV
- Pré/post charge
- Fonction contractile OG
- Volémie
- Valvulopathies

DTI anneau mitral



Après sérum PHY

Pas de modification significative de E' ni de E/E'



Après TNT

Relation ratio E/E' et PTDVG

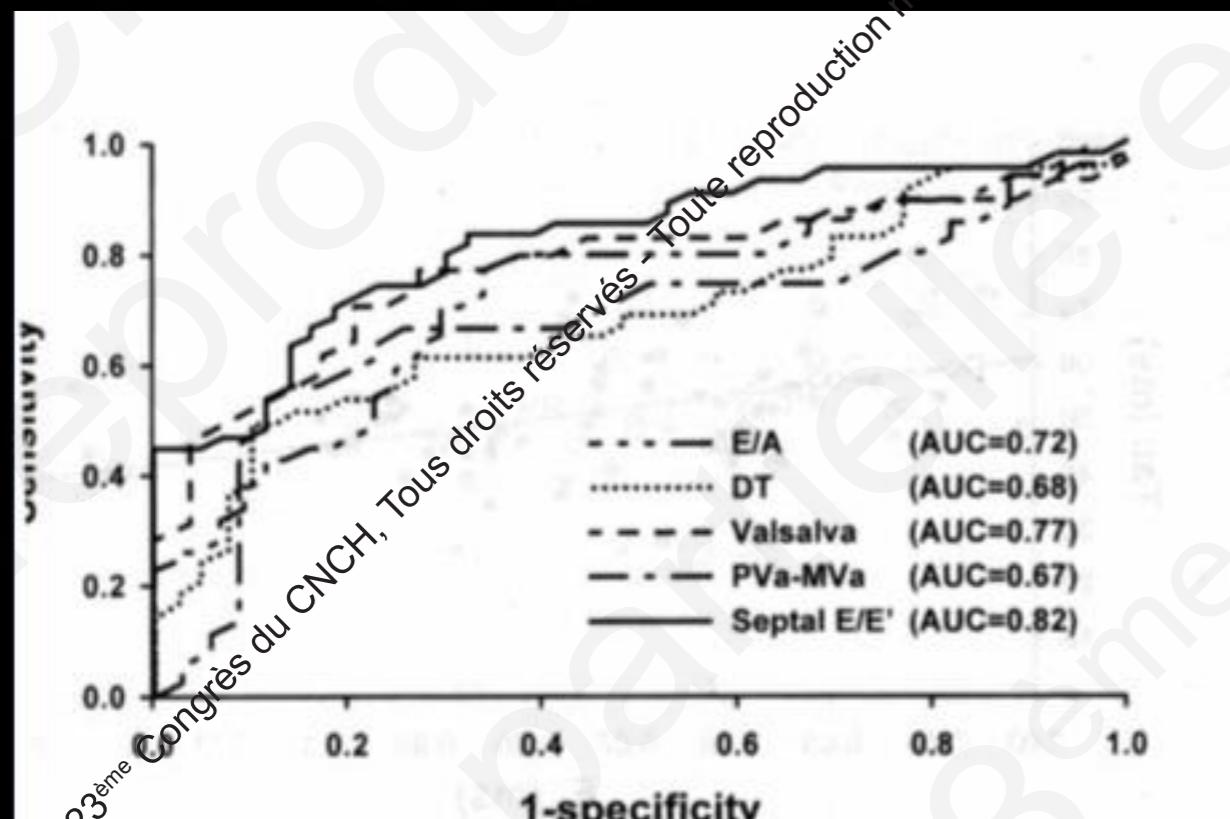


Figure 5. ROC curves for prediction of M-LVDP > 12 mm Hg for Doppler parameters. AUC indicates area under curve.

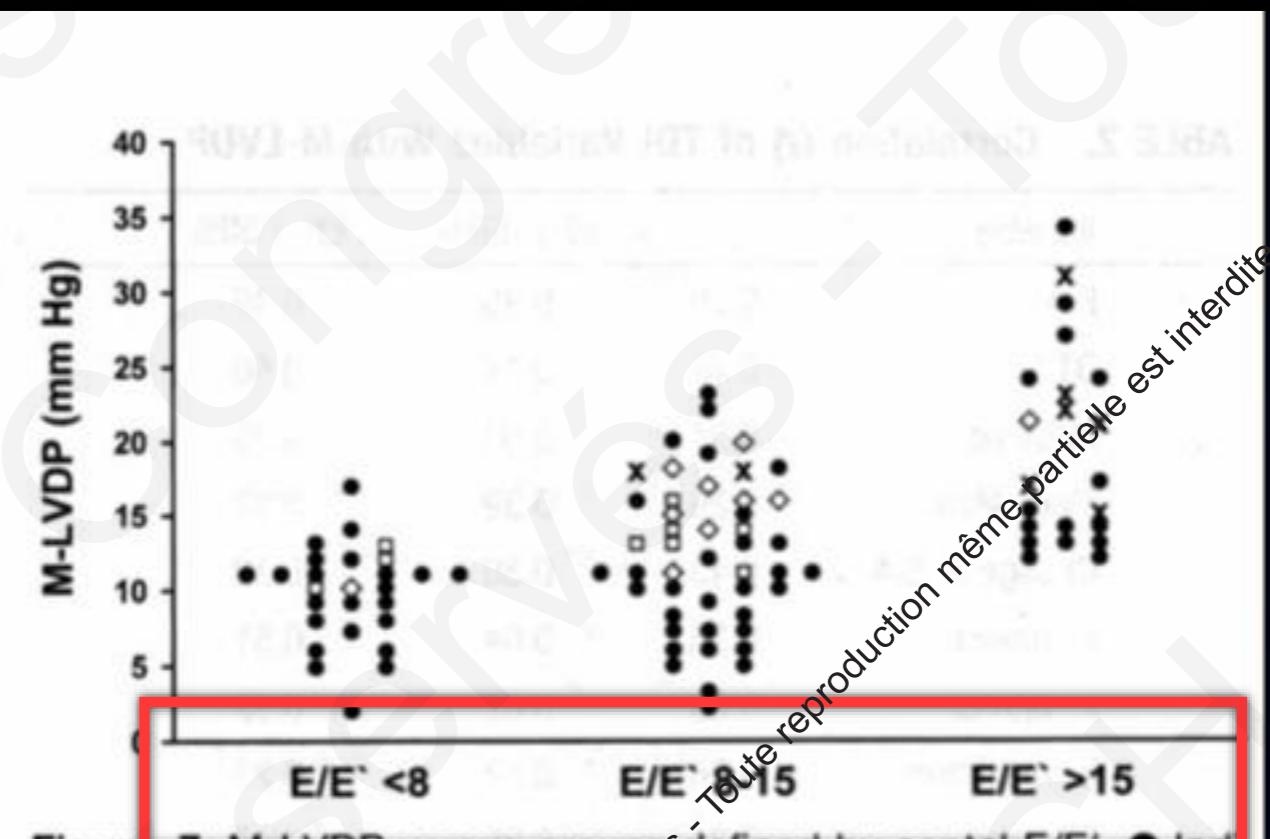


Figure 7. M-LVDP versus groups defined by septal E/E'. ● indicates patients without conventional Doppler variables suggesting increased filling pressure; □, patients with PVa 30 ms > MVa; ◇, patients with positive Valsalva maneuver; and X, > 15 .

85% de pts avec $E/E' < 8$ ont des PRVG basses et 100% des pts avec $E/E' > 15$ ont des PRVG hautes

Table 4 e' wave, E/e' ratio, left atrial volume (LA), and systolic pulmonary artery pressure (sPAP) according to age and gender

Parameters	20–40 years			40–60 years			≥60 years			Global cohort		
	Total (N = 172) n/ N (%)	Male (N = 79) n/ N (%)	Female (N = 93) n/N (%)	Total (N = 194) n/ N (%)	Male (N = 80) n/ N (%)	Female (N = 114) n/ N (%)	Total (N = 83) n/ N (%)	Male (N = 39) n/ N (%)	Female (N = 44) n/N (%)	Total (N = 449) n/ N (%)	Male (N = 198) n/ N (%)	Female (N = 251) n/N (%)
Septal e' <8 cm/s	2/170 (1.2)	2/79 (2.5)	0/91 (0)	38/193 (19.7)	13/80 (16.3)	25/113 (22.1)	46/83 (55.4)	22/39 (56.4)	24/44 (54.5)	86/446 (19.3)	37/198 (18.7)	49/248 (19.8)
Lateral e' <8 cm/s	2/167 (1.2)	1/76 (1.3)	1/91 (1.1)	11/192 (5.7)	4/80 (5.0)	7/112 (6.3)	15/79 (19.0)	5/36 (13.9)	10/43 (23.3)	28/438 (6.4)	10/192 (5.2)	18/246 (7.3)
Average sept-lat. E/e' >15	0/158 (0)	0/75 (0)	0/83 (0)	2/184 (1.1)	1/76 (1.3)	1/108 (0.9)	0/78 (0)	0/36 (0)	0/42 (0)	2/420 (0.5)	1/187 (0.5)	1/233 (0.4)
Lateral E/e' >13	0/159 (0)	0/75 (0)	0/84 (0)	3/184 (1.6)	1/76 (1.3)	2/108 (1.9)	0/78 (0)	0/36 (0)	0/42 (0)	3/421 (0.7)	1/187 (0.5)	2/234 (0.9)
LA volume >34 mL/m ^{2a}	18/149 (12.1)	9/68 (13.2)	9/81 (11.1)	30/169 (17.8)	15/68 (22.1)	15/101 (14.9)	10/66 (15.2)	5/33 (15.2)	5/33 (15.2)	58/384 (15.1)	29/169 (17.2)	29/215 (13.5)
LA volume >37 mL/m ^{2b}	10/149 (6.7)	6/68 (8.8)	4/81 (4.9)	18/169 (10.7)	8/68 (11.8)	10/101 (9.9)	6/66 (9.1)	3/33 (9.1)	3/33 (9.1)	34/384 (8.9)	17/169 (10.1)	17/215 (7.9)
LA volume >34 mL/m ^{2c}	4/124 (3.2)	2/58 (3.4)	2/66 (3.0)	19/131 (14.5)	7/54 (13.0)	12/77 (15.6)	5/53 (9.4)	4/26 (15.4)	1/27 (3.7)	28/308 (9.1)	13/138 (9.4)	15/170 (8.8)
LA volume >34 mL/m ^{2d}	10/117 (8.5)	4/53 (7.5)	6/64 (9.4)	18/127 (14.2)	7/51 (13.7)	11/76 (14.5)	3/50 (6.0)	2/24 (8.3)	1/26 (3.8)	31/294 (10.4)	13/128 (10.2)	18/166 (10.8)
LA volume >37 mL/m ^{2e}	5/117 (4.3)	2/53 (3.8)	3/64 (4.7)	10/127 (7.9)	5/51 (9.8)	5/76 (6.6)	1/50 (2.0)	0/24 (0.0)	1/26 (3.8)	16/294 (5.4)	7/128 (5.5)	9/166 (5.4)
LA volume >34 mL/m ^{2f}	20/117 (17.1)	8/53 (15.1)	12/64 (18.8)	33/127 (26.0)	16/51 (31.4)	17/76 (22.4)	8/50 (16.0)	6/24 (25.0)	2/26 (7.7)	61/294 (20.7)	30/128 (23.4)	31/166 (18.7)
LA volume >37 mL/m ^{2g}	12/117 (10.3)	6/53 (11.3)	6/64 (9.4)	21/127 (16.5)	11/51 (21.6)	10/76 (13.2)	4/50 (8.0)	3/24 (12.5)	1/26 (3.8)	37/294 (12.6)	20/128 (15.6)	17/166 (10.2)
sPAP >36 mmHg	1/106 (0.9)	1/48 (2.1)	0/58 (0.0)	0/131 (0.0)	0/57 (0.0)	0/74 (0.0)	0/57 (0.0)	0/24 (0.0)	0/33 (0.0)	1/294 (0.3)	1/129 (0.8)	0/165 (0.0)
sPAP >45 mmHg	0/106 (0.0)	0/48 (0.0)	0/58 (0.0)	0/131 (0.0)	0/57 (0.0)	0/74 (0.0)	0/57 (0.0)	0/24 (0.0)	0/33 (0.0)	0/294 (0.0)	0/129 (0.0)	0/165 (0.0)

^aLA volume >34 mL/m² by single plane area-length method (four-chamber view).

^bLA volume >37 mL/m² by single plane area-length method (four-chamber view).

^cLA volume >34 mL/m² by single plane Simpson method (four-chamber view).

^dLA volume >34 mL/m² by biplane Simpson method.

^eLA volume >37 mL/m² by biplane Simpson method.

^fLA volume >34 mL/m² by biplane area-length method.

^gLA volume >34 mL/m² by biplane area-length method.

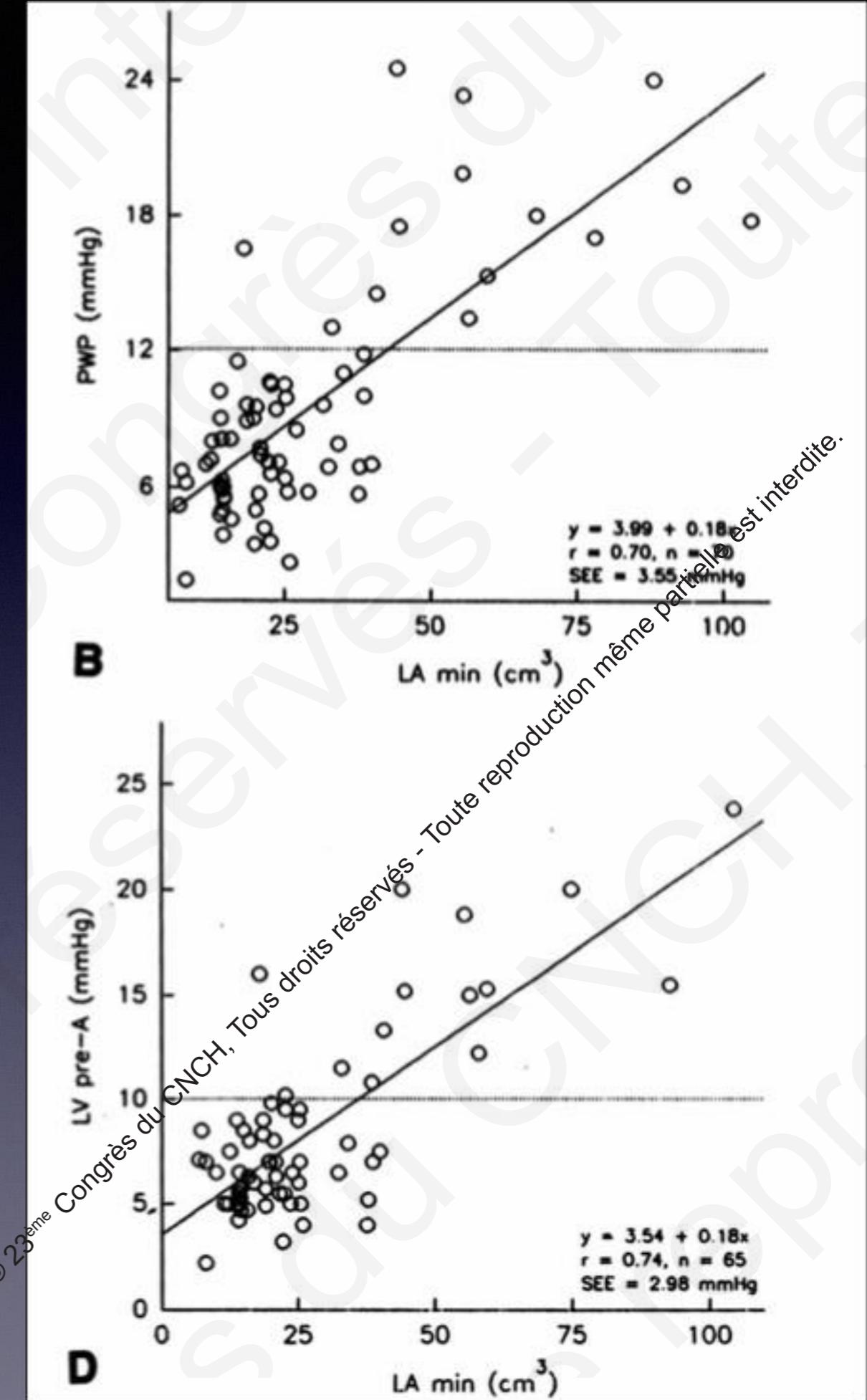
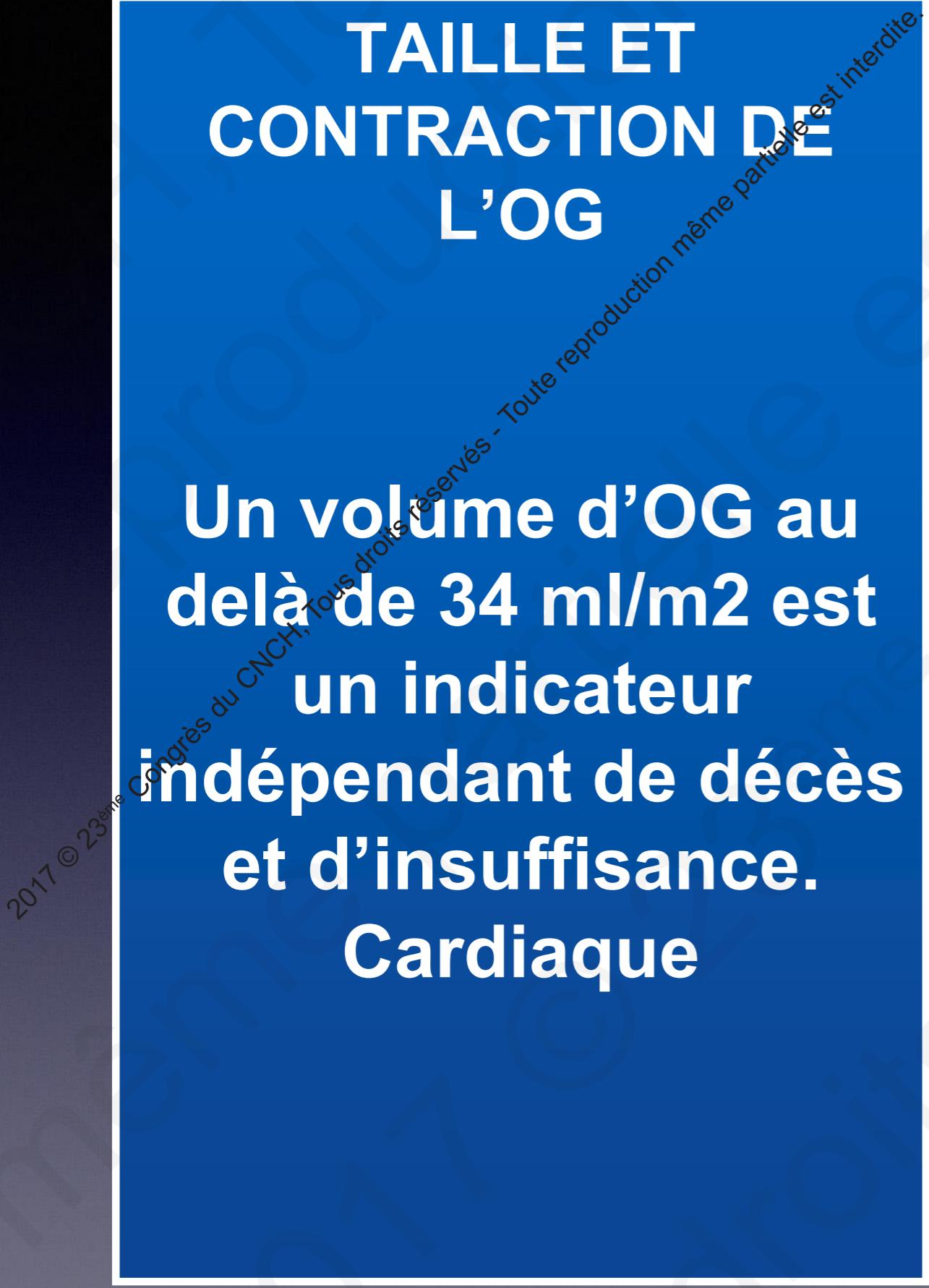
Autres paramètres

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TAILLE ET CONTRACTION DE L'OG

Un volume d'OG au delà de 34 ml/m² est un indicateur indépendant de décès et d'insuffisance. Cardiaque



PAPs et dysfonction diastolique

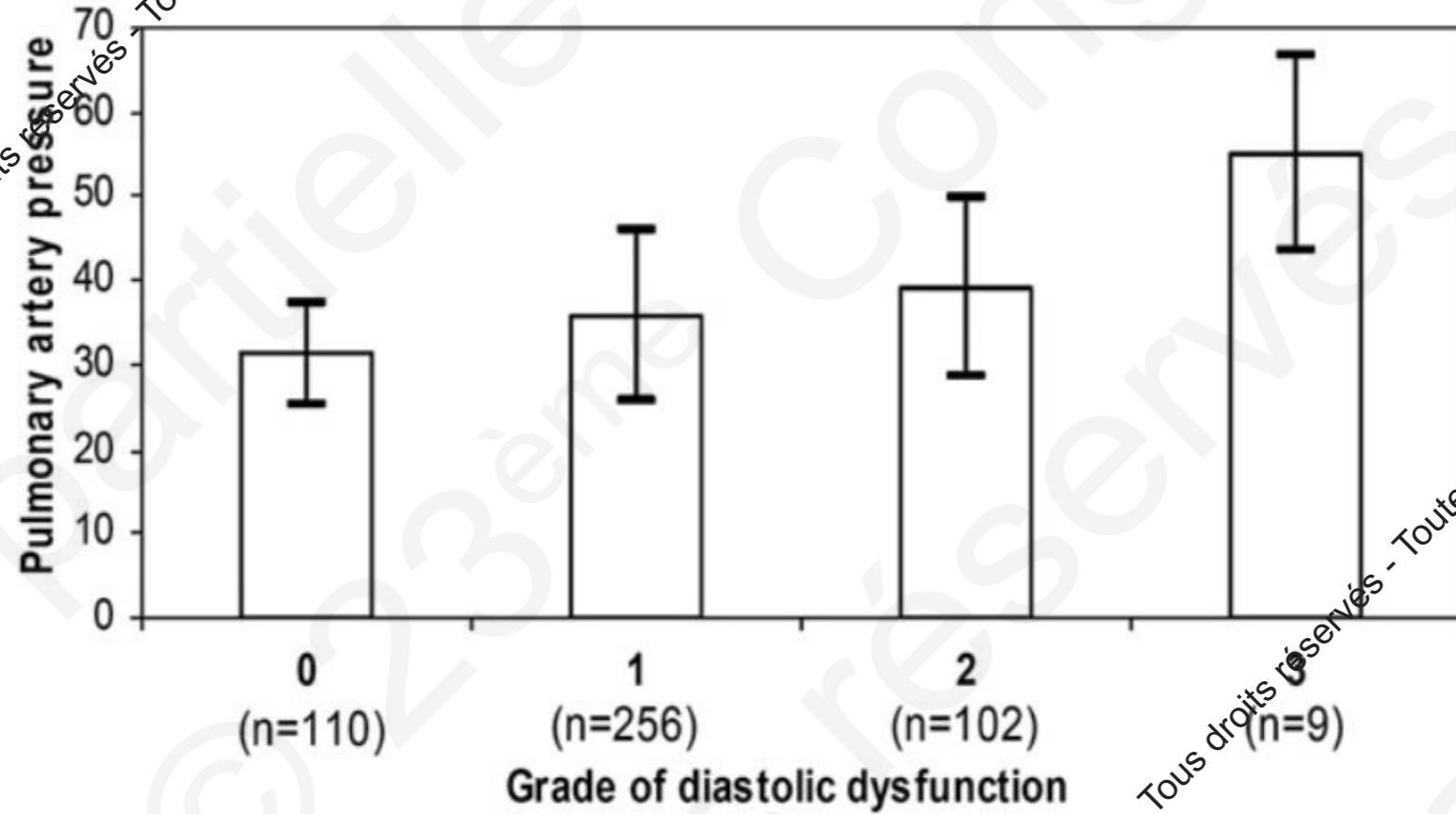


Fig. 1. Mean pulmonary artery pressure (mm Hg) by grade of diastolic dysfunction. $P < 0.001$ by one-way ANOVA for the differences was between each 2 groups of diastolic dysfunction.

PAPs et dysfonction diastolique

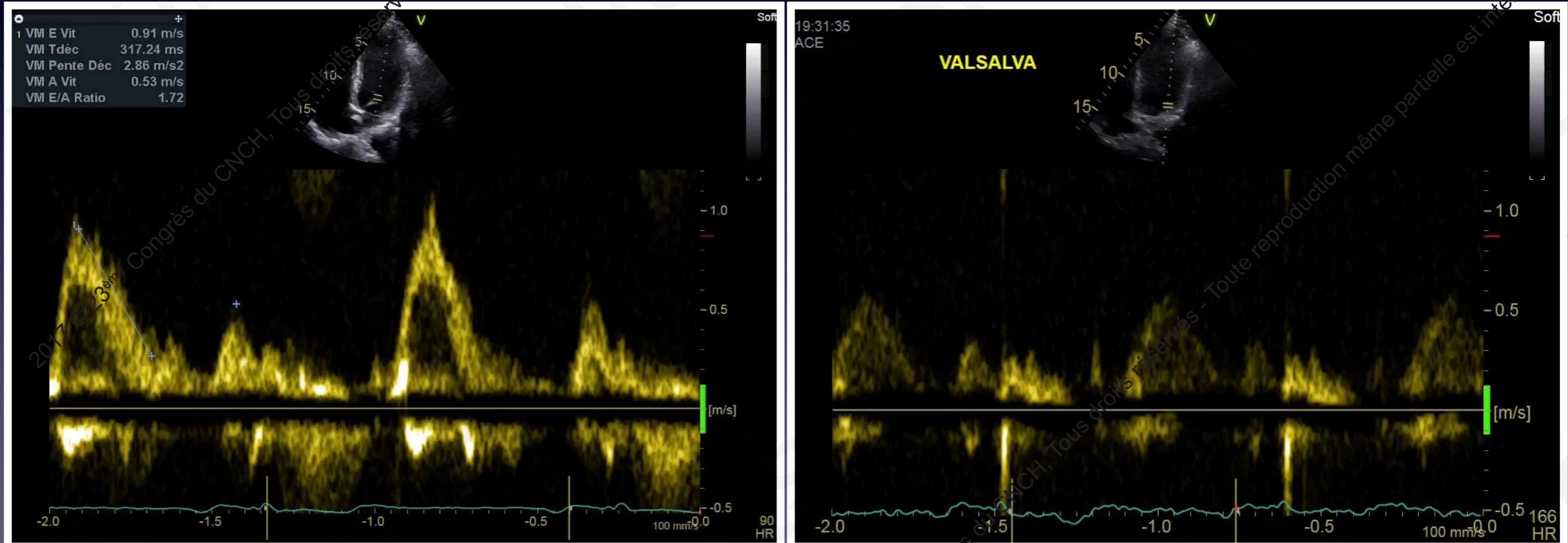
Table 3 Proportion of normal subjects with abnormal LA volume index and Doppler velocities

Parameter	20–40 y			40–60 y			≥60 y			Global cohort		
	Total (N = 172) n/N (%)	Male (N = 79) n/N (%)	Female (N = 93) n/N (%)	Total (N = 194) n/N (%)	Male (N = 80) n/N (%)	Female (N = 114) n/N (%)	Total (N = 83) n/N (%)	Male (N = 39) n/N (%)	Female (N = 44) n/N (%)	Total (N = 449) n/N (%)	Male (N = 198) n/N (%)	Female (N = 251) n/N (%)
Septal e' < 8 cm/sec	2/170 (1.2)	2/79 (2.5)	0/91 (0)	38/193 (19.7)	13/80 (16.3)	25/113 (22.1)	46/83 (55.4)	22/39 (56.4)	24/44 (54.5)	86/446 (19.3)	37/198 (18.7)	49/248 (19.8)
Lateral e' < 8 cm/sec	2/167 (1.2)	1/76 (1.3)	1/91 (1.1)	11/192 (5.7)	4/80 (5.0)	7/112 (6.3)	15/79 (19.0)	5/36 (13.9)	10/43 (23.3)	28/438 (6.4)	10/192 (5.2)	18/246 (7.3)
Lateral e' < 10 cm/sec	3/167 (1.8)	2/76 (2.6)	1/91 (1.1)	30/192 (15.6)	9/80 (11.3)	21/112 (18.8)	41/79 (51.9)	17/36 (47.2)	24/43 (55.8)	74/438 (16.9)	28/192 (14.6)	46/246 (18.7)
Average (septal-lateral) E/e' > 14	0/158 (0)	0/75 (0)	0/83 (0)	3/184 (1.6)	1/76 (1.3)	2/108 (1.9)	1/78 (1.3)	1/36 (2.8)	0/42 (0)	4/420 (1.0)	2/187 (1.1)	2/233 (0.9)
Septal E/e' > 15	0/162 (0)	0/78 (0)	0/84 (0)	2/185 (1.1)	1/76 (1.3)	1/109 (0.9)	3/81 (3.7)	2/38 (5.3)	1/43 (2.3)	5/428 (1.2)	3/192 (1.6)	2/236 (0.8)
Lateral E/e' > 13	0/159 (0)	0/75 (0)	0/84 (0)	3/184 (1.6)	1/76 (1.3)	2/108 (1.9)	0/78 (0)	0/36 (0)	0/42 (0)	3/421 (0.7)	1/187 (0.5)	2/234 (0.9)
LA volume index > 34 mL/m ² (*)	10/117 (8.5)	4/53 (7.5)	6/64 (9.4)	18/127 (14.2)	7/51 (13.7)	11/76 (14.5)	3/50 (6.0)	2/24 (8.3)	1/26 (3.8)	31/294 (10.5)	13/128 (10.2)	18/166 (10.8)
SPAP > 36 mm Hg	1/106 (0.9)	1/48 (2.1)	0/58 (0.0)	0/131 (0.0)	0/57 (0.0)	0/74 (0.0)	0/57 (0.0)	0/24 (0.0)	0/33 (0.0)	1/294 (0.3)	1/129 (0.8)	0/165 (0.0)
SPAP > 45 mm Hg	0/106 (0.0)	0/48 (0.0)	0/58 (0.0)	0/131 (0.0)	0/57 (0.0)	0/74 (0.0)	0/57 (0.0)	0/24 (0.0)	0/33 (0.0)	0/294 (0.0)	0/129 (0.0)	0/165 (0.0)

SPAP, Systolic pulmonary artery pressure.

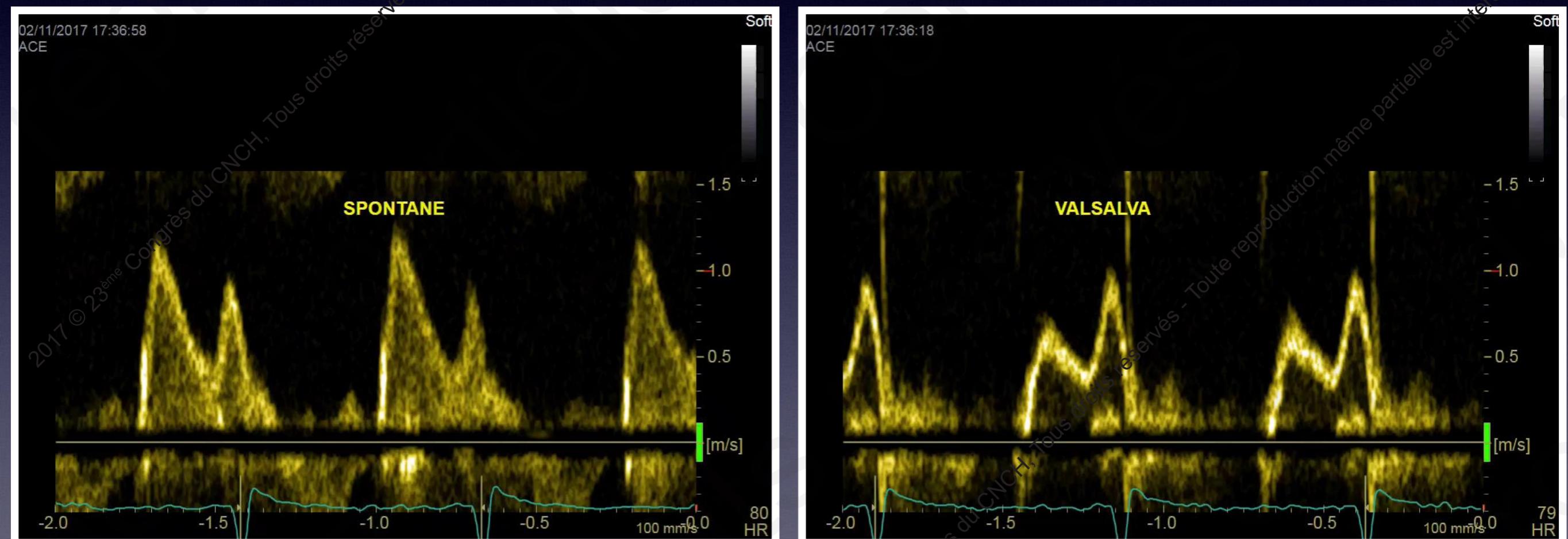
*LA volume index > 34 mL/m² by biplane Simpson method (adapted from Caballero et al.⁵²).

Valsalva



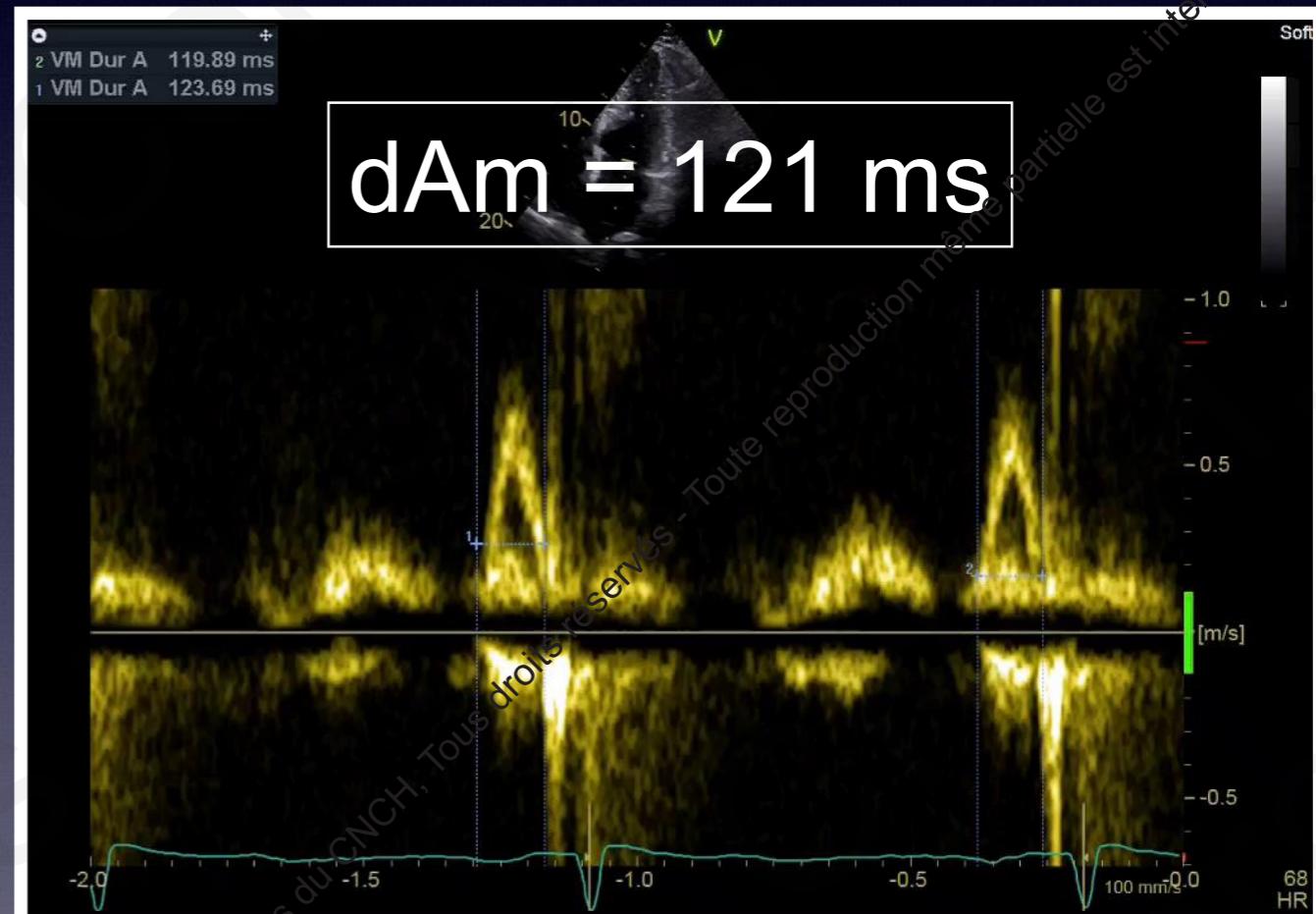
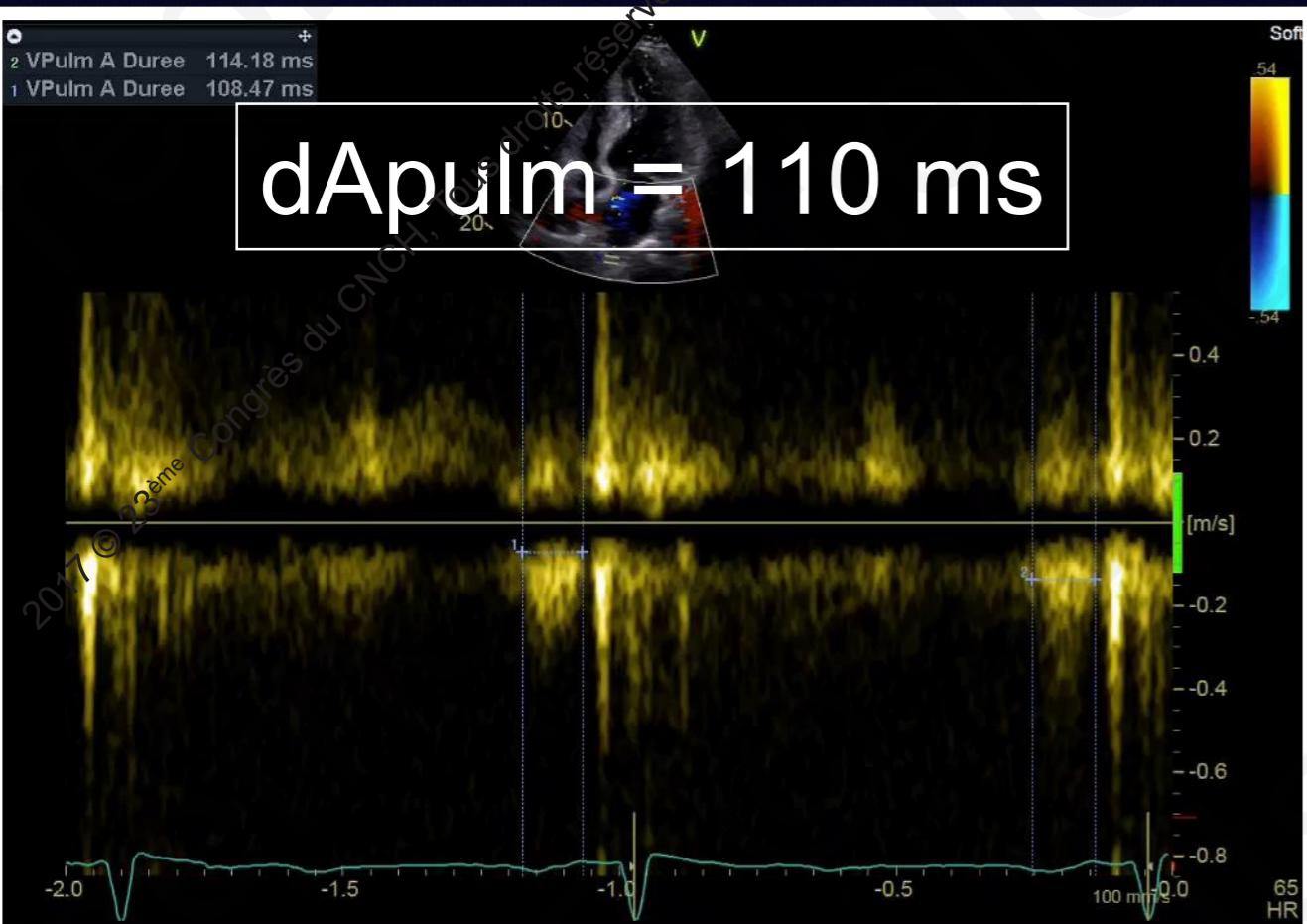
Sujet normal
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Valsalva



PRVG augmentées

dAp-dAm



PRVG normales

Méthodes échographiques d'évaluation de la fonction diastolique

Table 1 (Continued)^a

Variable	Acquisition	Analysis
PV S/D ratio	See above for acquisition of pulmonary vein S and D velocities. PV S wave divided by D-wave velocity or PV S wave time-velocity integral/PV D wave time-velocity integral.	
CW Doppler: TR systolic jet velocity (m/sec)	1. Parasternal and apical four-chamber view with color flow imaging to obtain highest Doppler velocity aligned with CW. 2. Adjust gain and contrast to display complete spectral envelope without signal spikes or feathering	Peak modal velocity during systole at leading edge of spectral waveform
Secondary measures		(Continued)
Color M-mode Vp (cm/sec)	Apical four-chamber with color flow imaging for M-mode cursor position, shift color baseline in direction of mitral valve inflow to lower velocity scale for red/yellow inflow velocity profile	Slope of inflow from MV plane into LV chamber during early diastole (cm/sec/cm distance)
IVRT	Apical long-axis or five-chamber view, using CW Doppler and placing sample volume in LV outflow tract to simultaneously display end of aortic ejection and onset of mitral inflow.	Time between aortic valve closure and MV opening. For IVRT, sweep speed should be 100 mm/sec.
TE-e'	Apical four-chamber view with proper alignment to acquire mitral inflow at mitral valve tips and using tissue Doppler to acquire septal and lateral mitral annular velocities.	Time interval between peak of R wave in QRS complex and onset of mitral E velocity is subtracted from time interval between QRS complex and onset of e' velocity. RR intervals should be matched and gain and filter settings should be optimized to avoid high gain and filter settings. For time intervals, sweep speed should be 100 mm/sec.

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Variable	Acquisition	Analysis
Peak E-wave velocity (cm/sec)	1. Apical four-chamber with color flow imaging for optimal alignment of PW Doppler with blood flow. 2. PW Doppler sample volume (1–3 mm axial size) between mitral leaflet tips. 3. Use low wall filter setting (100–200 MHz) and low signal gain. 4. Optimal spectral waveforms should not display spikes or feathering.	Peak modal velocity in early diastole (after ECG T wave) at the leading edge of spectral waveform
Peak A-wave velocity (cm/sec)	1. Apical four-chamber with color flow imaging for optimal alignment of PW Doppler with blood flow 2. PW Doppler sample volume (1–3 mm axial size) between mitral leaflet tips. 3. Use low wall filter setting (100–200 MHz) and low signal gain. 4. Optimal spectral waveforms should not display spikes or feathering.	Peak modal velocity in late diastole (after ECG P wave) at the leading edge of spectral waveform
MV A duration (msec)	1. Apical four-chamber with color flow imaging for optimal alignment of PW Doppler with blood flow. 2. PW Doppler sample volume (1–3 mm axial size) at level of mitral annulus (limited data on how duration compares between annulus and leaflet tips) 3. Use low wall filter setting (100–200 MHz) and low signal gain. 4. Optimal spectral waveforms should not display spikes or feathering.	Time interval from A-wave onset to end of A wave at zero baseline. If E and A are fused (E velocity > 20 cm/sec when A velocity starts), A-wave duration will often be longer because of increased atrial filling stroke volume.
MV E/A ratio	See above for proper technique of acquisition of E and A velocities.	MV E velocity divided by A-wave velocity
MV DT (msec)	Apical four-chamber: pulsed Doppler sample volume between mitral leaflet tips	Time interval from peak E-wave along the slope of LV filling extrapolated to the zero-velocity baseline.
Pulsed-wave TDI e' velocity (cm/sec)	1. Apical four-chamber view: PW Doppler sample volume (usually 5–10 mm axial size) at lateral and septal basal regions so average e' velocity can be computed. 2. Use ultrasound system presets for wall filter and lowest signal gain. 3. Optimal spectral waveforms should not display spikes or feathering.	Peak modal velocity in early diastole at the leading edge of spectral waveform

Variable	Acquisition	Analysis
Peak E-wave velocity (cm/sec)	1. Apical four-chamber with color flow imaging for optimal alignment of PW Doppler with blood flow. 2. PW Doppler sample volume (1–3 mm axial size) between mitral leaflet tips. 3. Use low wall filter setting (100–200 MHz) and low signal gain. 4. Optimal spectral waveforms should not display spikes or feathering.	Peak modal velocity in early diastole (after ECG T wave) at the leading edge of spectral waveform
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Secondary measures		(Continued)
Color M-mode Vp (cm/sec)	Apical four-chamber with color flow imaging for M-mode cursor position, shift color baseline in direction of mitral valve inflow to lower velocity scale for red/yellow inflow velocity profile	Slope of inflow from MV plane into LV chamber during early diastole (cm/sec/cm distance)
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ASE/EACVI GUIDELINES AND STANDARDS

Recommendations for the Evaluation of Left Ventricular Diastolic Function by Echocardiography: An Update from the American Society of Echocardiography and the European Association of Cardiovascular Imaging

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Frank A. Flachskampf, MD, PhD, FESC,² Thierry C. Gillebert, MD, PhD, FESC,² Allan L. Klein, MD, FASE,¹
Patrizio Lancellotti, MD, PhD, FESC,² Paolo Marino, MD, FESC,² Jae K. Oh, MD,¹
Bogdan Alexandru Popescu, MD, PhD, FESC, FASE,² and Alan D. Waggoner, MHS, RDCS¹, Houston, Texas;
Oslo, Norway; Phoenix, Arizona; Nashville, Tennessee; Hamilton, Ontario, Canada; Uppsala, Sweden; Ghent and Liège, Belgium; Cleveland, Ohio; Novara, Italy; Rochester, Minnesota; Bucharest, Romania; and St. Louis, Missouri

(J Am Soc Echocardiogr 2016;29:277-314.)

Algorithme FE préservée

A

- 1-Average E/e' > 14**
- 2-Septal e' velocity < 7 cm/s or Lateral e' velocity <10 cm/s**
- 3-TR velocity > 2.8 m/s**
- 4-LA volume index >34ml/m²**

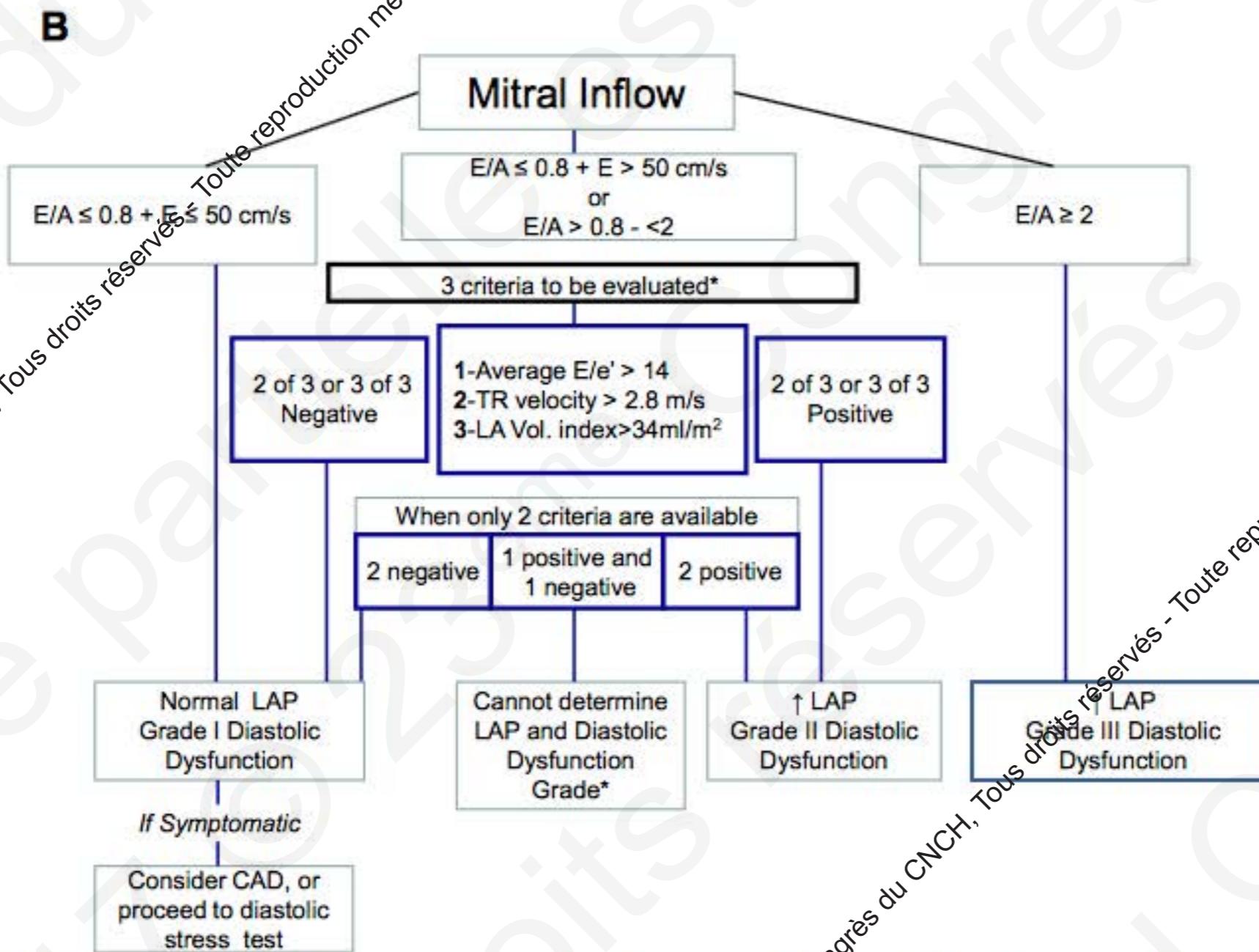
Normal Diastolic function

Indeterminate

Diastolic Dysfunction

6
ive

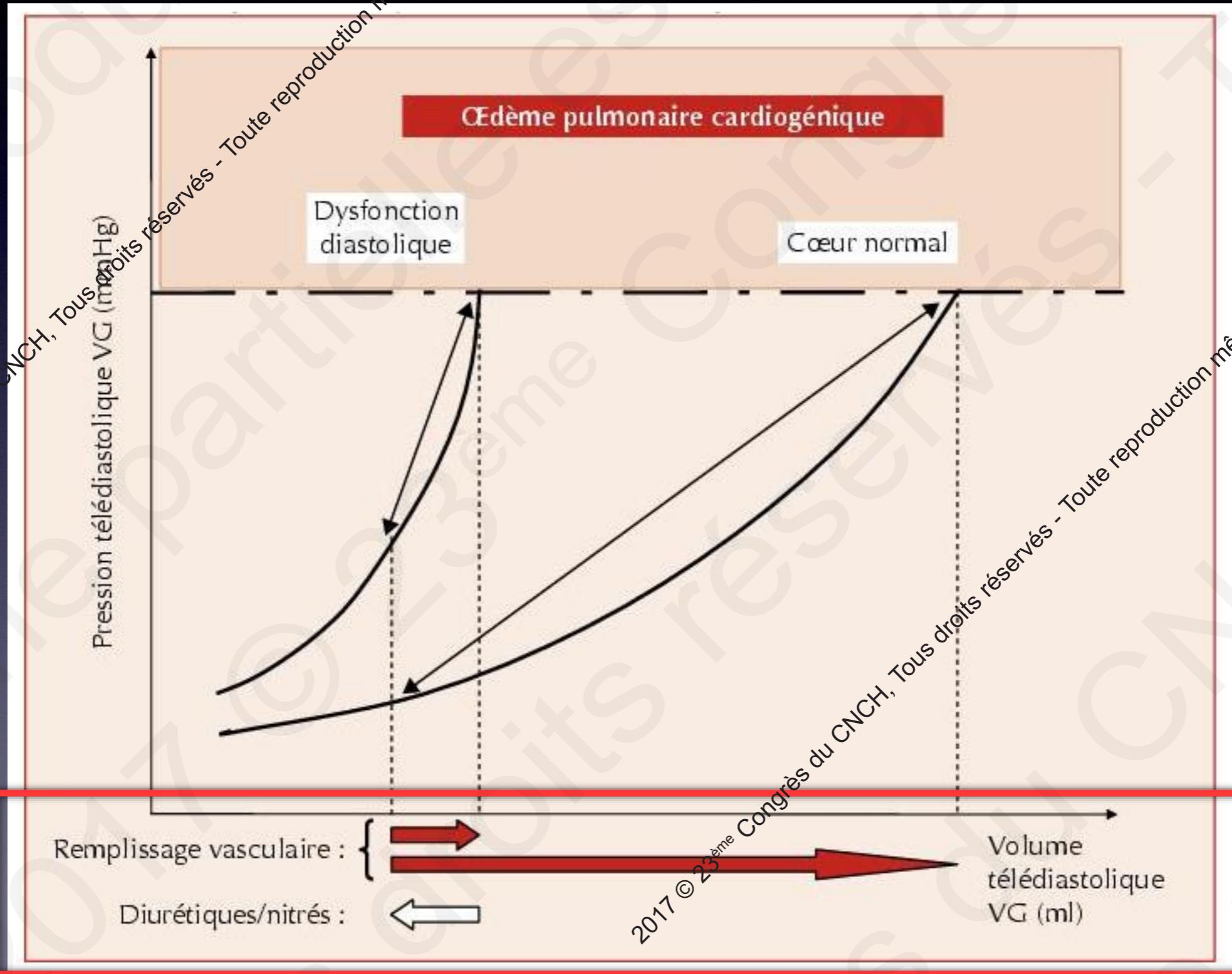
Algorithme FE altérée et indéterminé

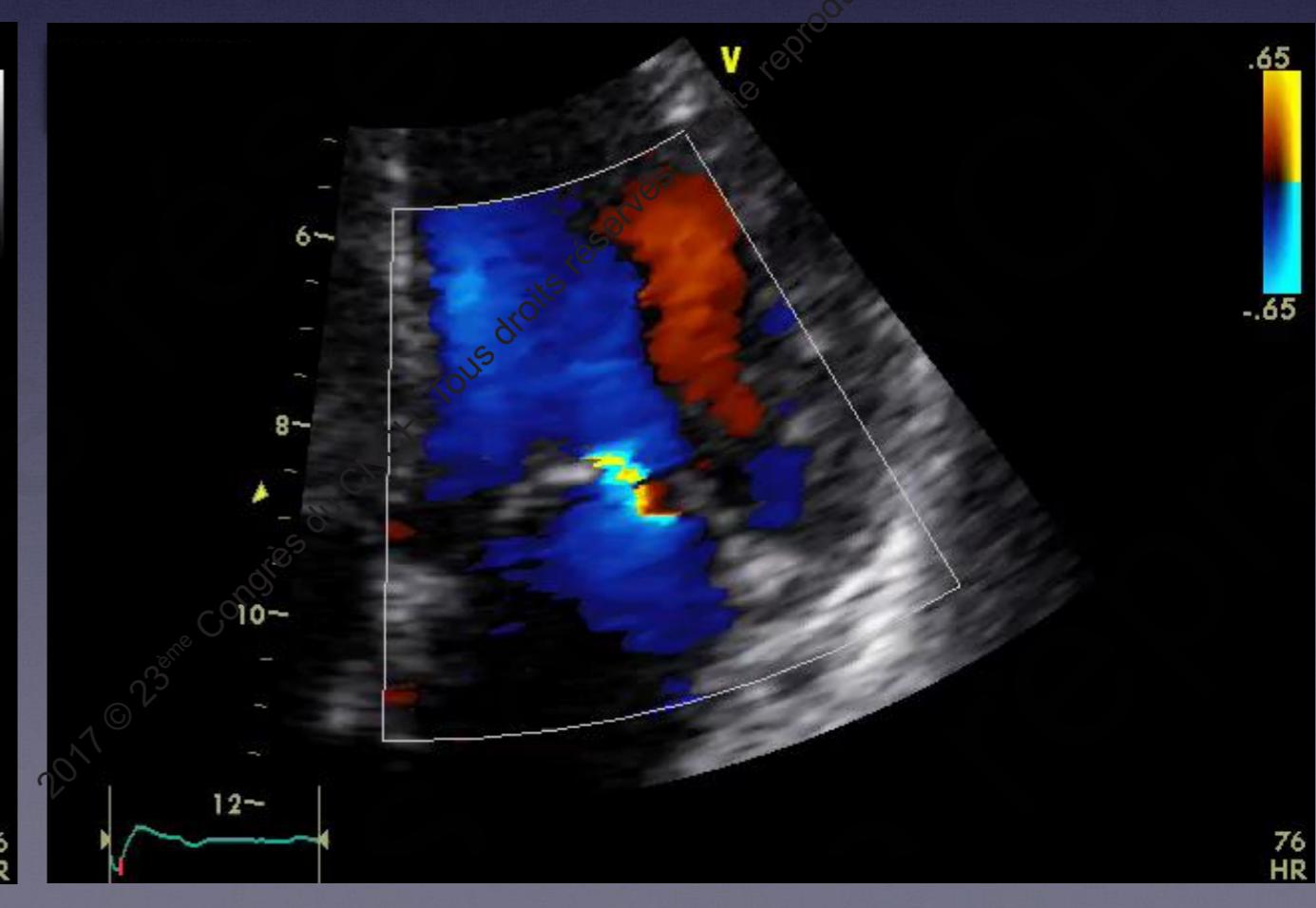
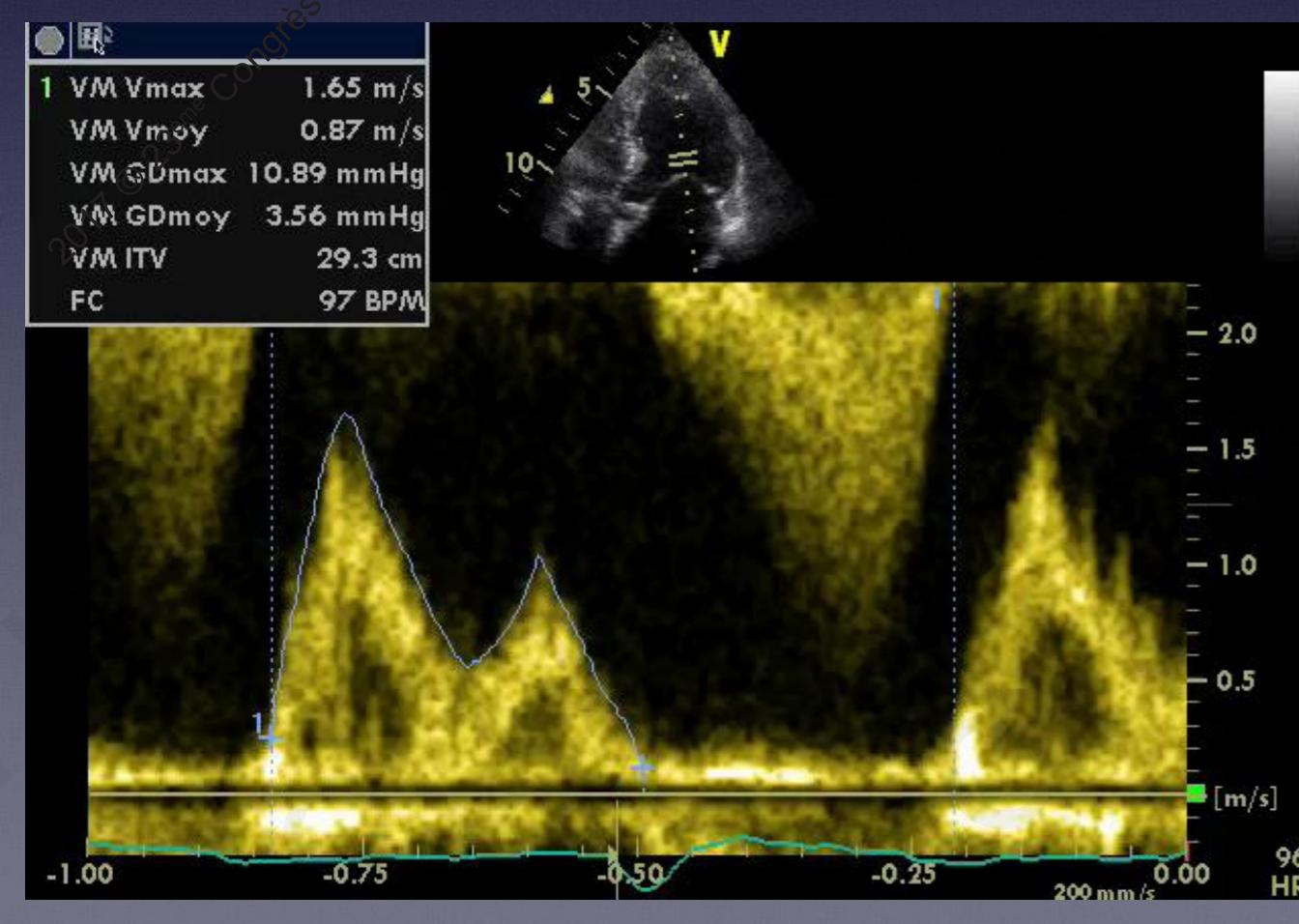
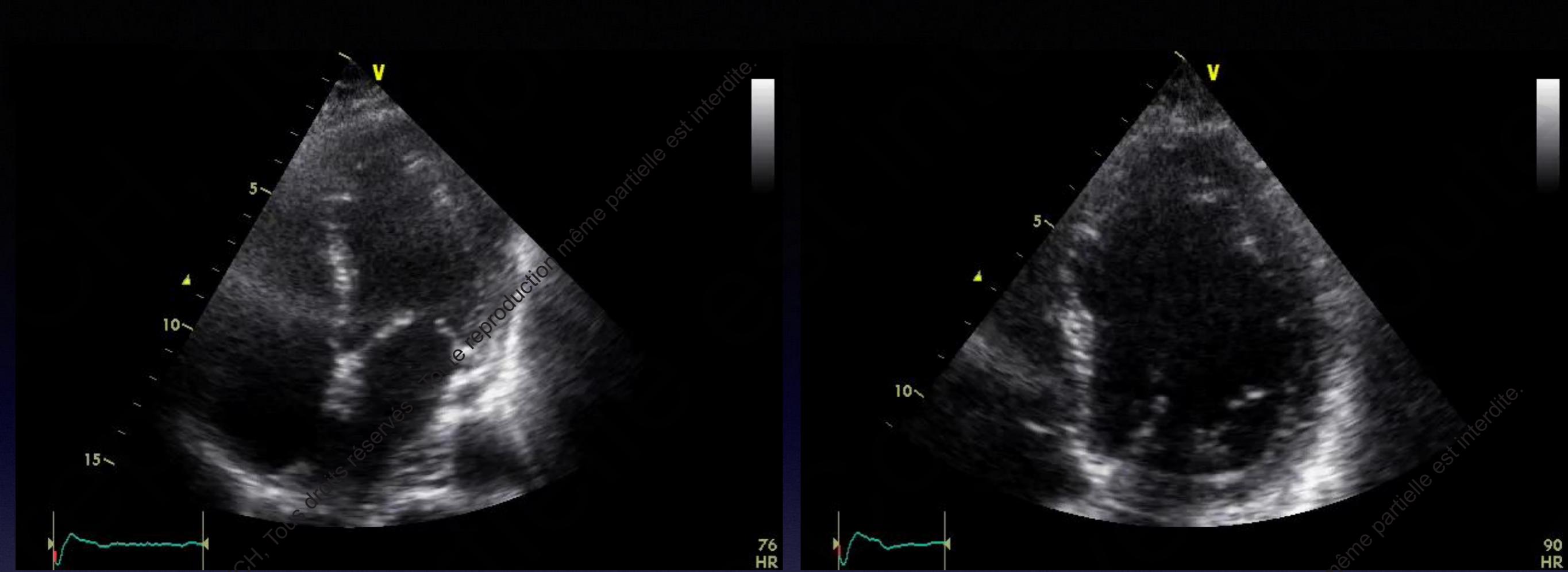


(* : LAP indeterminate if only 1 of 3 parameters available. Pulmonary vein S/D ratio <1 applicable to conclude elevated LAP in patients with depressed LV EF)

Figure 8 (A) Algorithm for diagnosis of LV diastolic dysfunction in subjects with normal LVEF. **(B)** Algorithm for estimation of LV filling pressures and grading LV diastolic function in patients with depressed LVEFs and patients with myocardial disease and normal LVEF after consideration of clinical and other 2D data.

Ne pas attendre l'arrivée des troupes fraîches



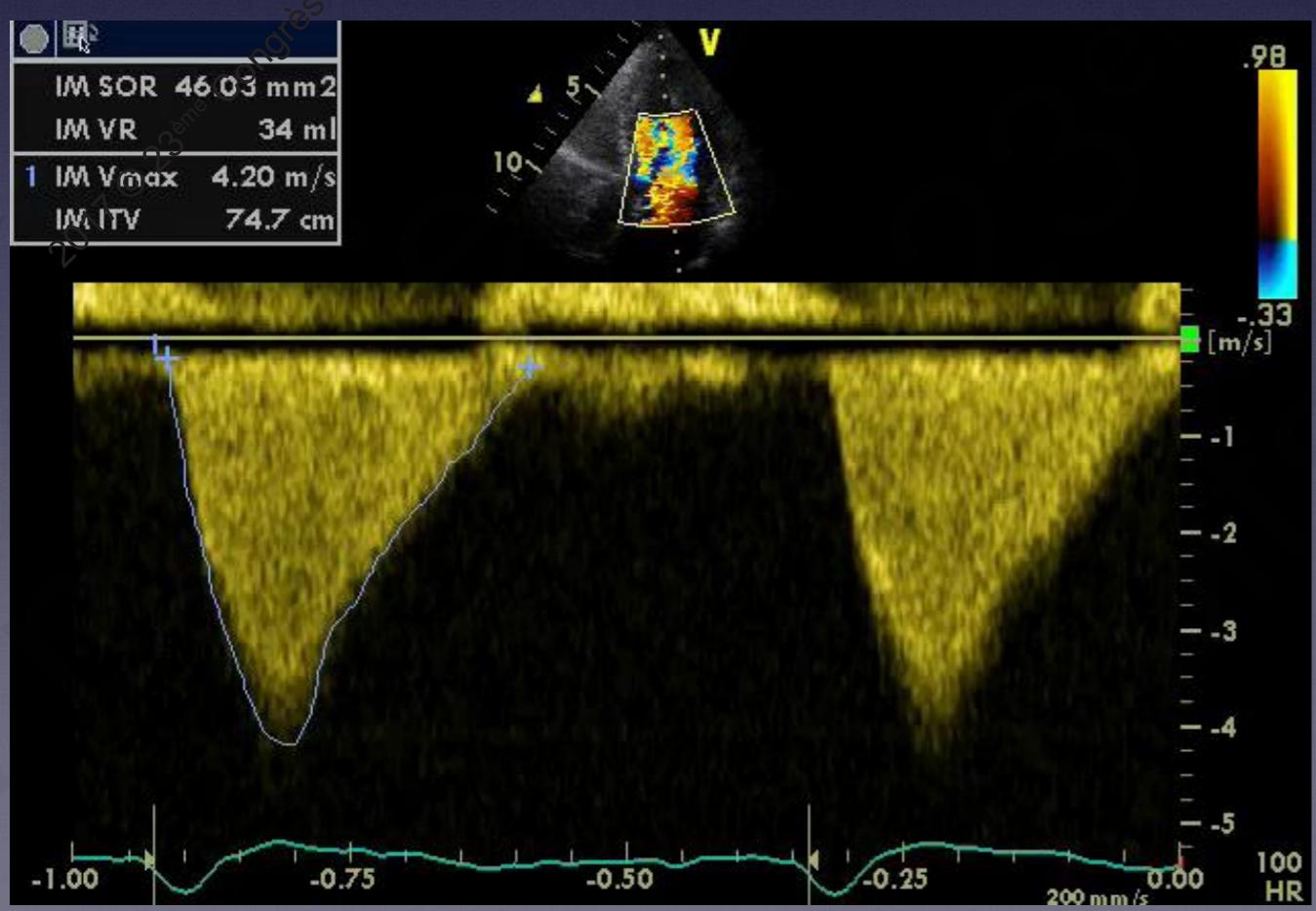




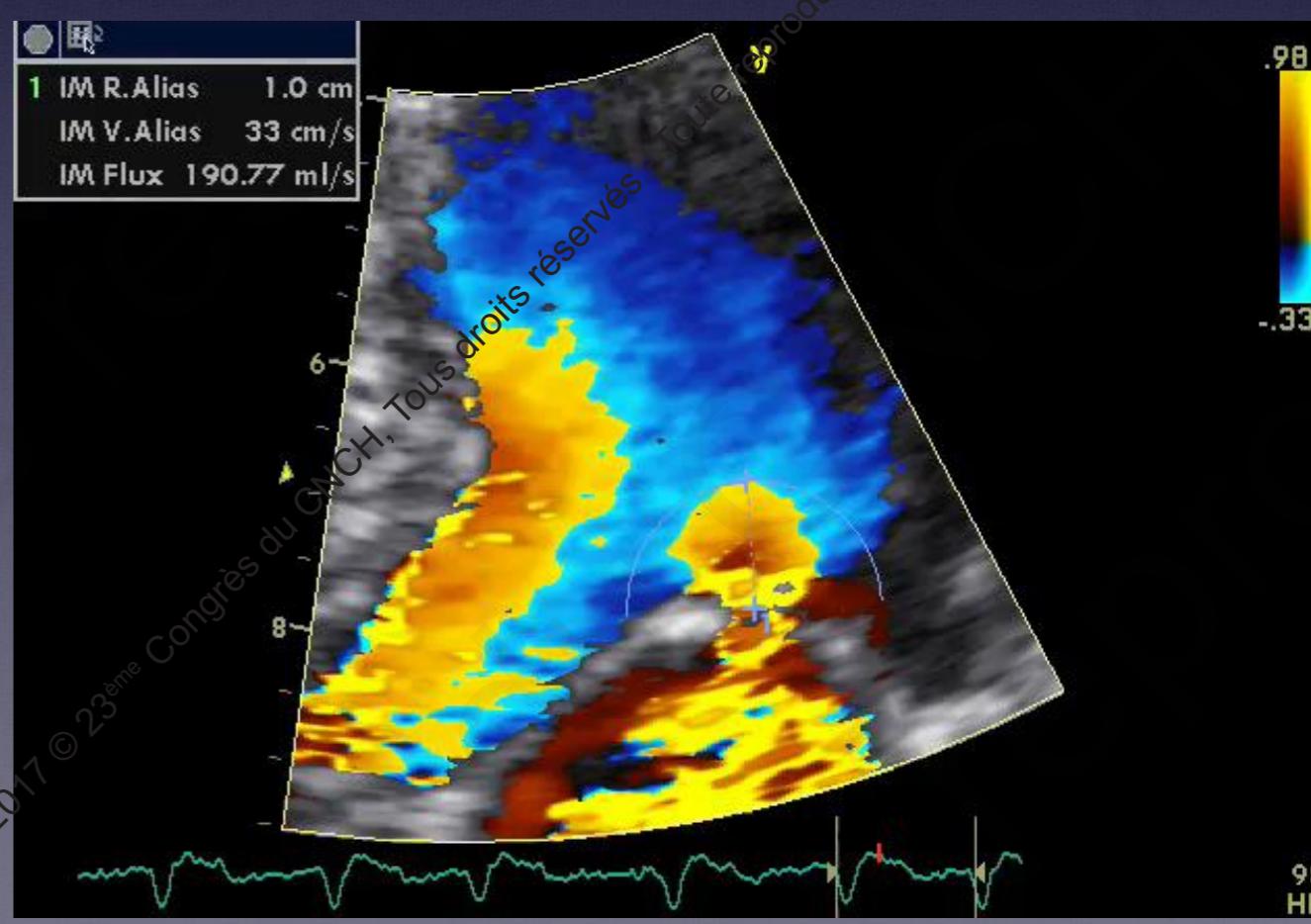
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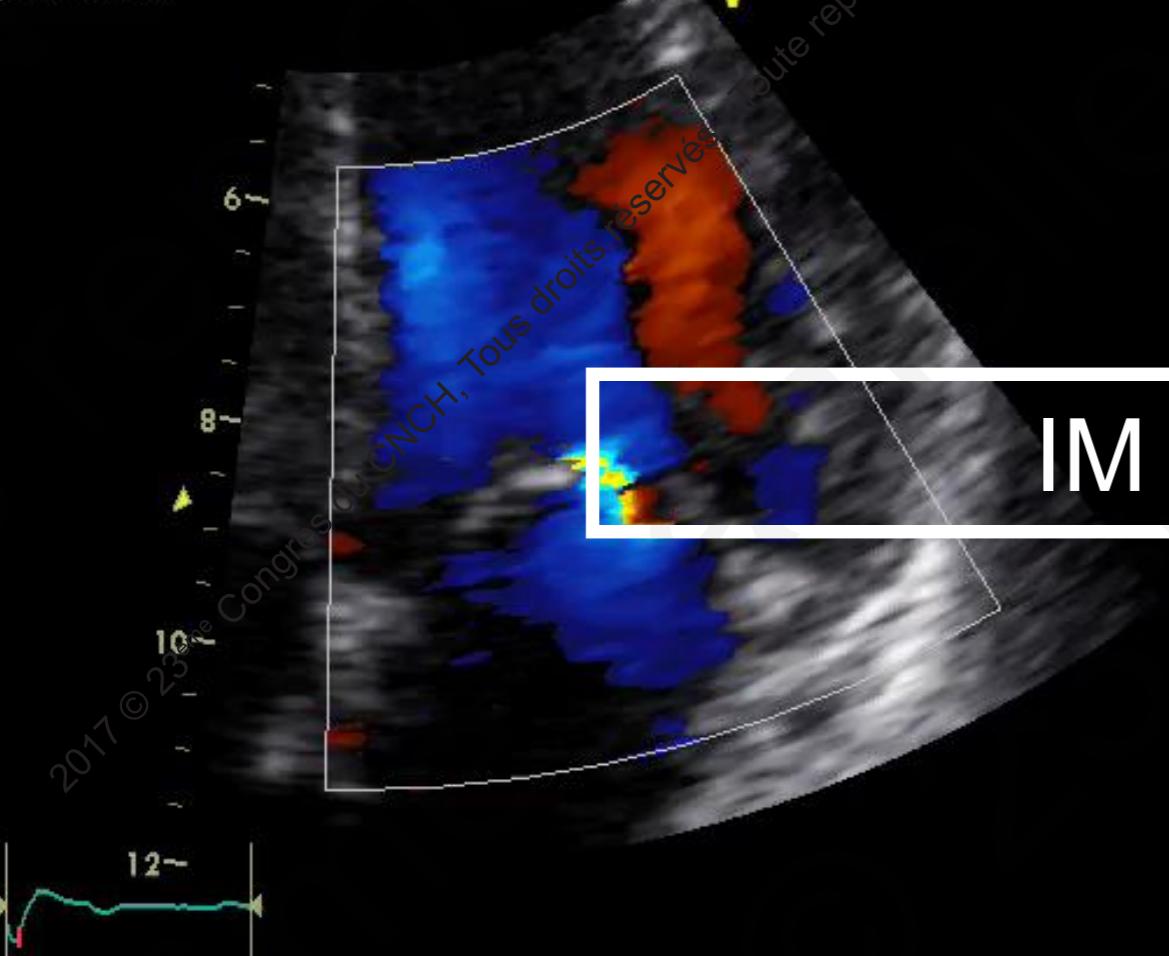
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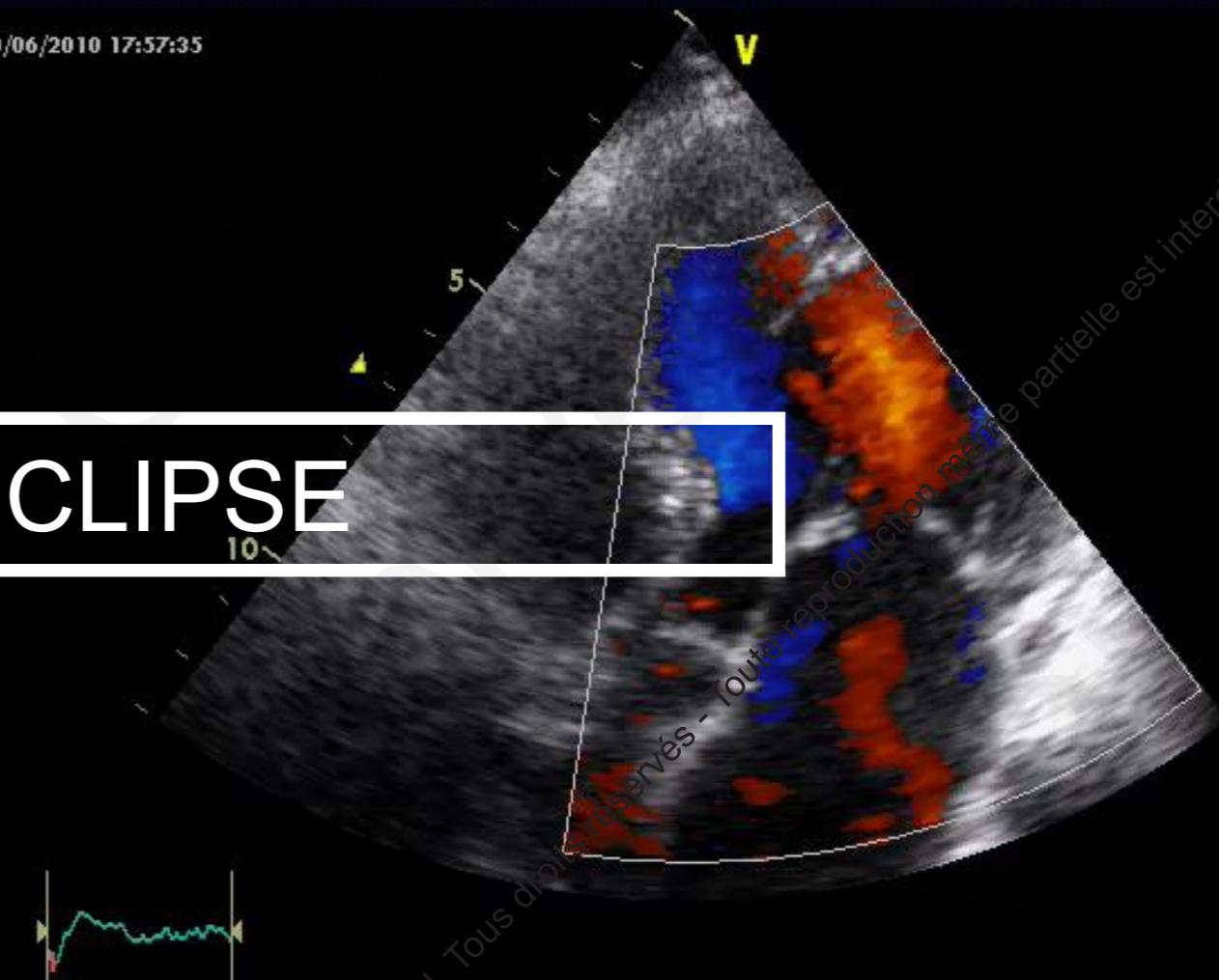
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The use of echocardiography in acute cardiovascular care: Recommendations of the European Association of Cardiovascular Imaging and the Acute Cardiovascular Care Association

Patrizio Lancellotti^{1*}, Susanna Price^{2*}, Thor Edvardsen³, Bernard Cosyns⁴, Aleksandar N. Neskovic⁵, Raluca Dulgheru¹, Frank A. Flachskampf⁶, Christian Hassager⁷, Agnes Pasquet⁸, Luna Gargani⁹, Maurizio Galderisi¹⁰, Nuno Cardim¹¹, Kristina H. Haugaa³, Arnaud Ancion¹, Jose-Luis Zamorano¹², Erwan Donal¹³, Héctor Bueno¹⁴, and Gilbert Habib¹⁵

Table 2 Echocardiographic signs indicative or suggestive of the cause of clinical admission in acute cardiovascular conditions

Systolic heart failure	Heart failure with preserved left ventricular ejection	Pulmonary embolism	Tamponade ^e
(1) LVEF<45–50% ^a (2) LVEDD >55 mm and/or >32 mm/m ² (3) LVESD >45 mm ^a and/or 25 mm/m ² (4) LVEDV>97 mL/m ² (5) LVESV>43 mL/m ² (6) Abnormal wall motion (7) Functional MR and/or TR (8) Peak tricuspid velocity >3 m/s (9) Aortic time velocity integral <15 cm ^a (10) Diastolic dysfunction (E/A ≥ 2 + DT<150 ms indicates increased LV filling pressures) ^b (11) Ultrasound lung comets ^c	(1) LVEF ≥ 50% ^a (2) LVEDV<97 mL/m ² (3) LVESV<43 mL/m ² (4) E–e' ≥ 13 ^b (5) Ar – A ≥ 30 ms (6) LA volume ≥ 34 mL/m ² (7) Peak tricuspid velocity>3 m/s (8) Ultrasound lung comets ^c + signs and symptoms of heart failure	(1) Thrombus into right chambers (2) Abnormal septal motion (3) Dilatation RA, RV (end-diastolic RV/LV diameter>0.6 or area>1.0) (4) Global RV hypokinesia (5) McConnell sign hyperkinesia (6) Mild to severe TR (7) Pulmonary hypertension around 40–50 mmHg (>60 mmHg in the case of pre-existing pulmonary hypertension)	(1) Usually large pericardial effusion (2) Swinging heart (3) RA collapse (rarely LA) (4) Diastolic collapse of the anterior RV-free wall (rarely LV) (5) IVC dilatation (no collapse with inspiration) (6) TV flow increases and MV flow decreases during inspiration (reverse in expiration) (7) Systolic and diastolic flows are reduced in systemic veins in expiration and reverse flow with atrial contraction is increased



EN CONCLUSION

- ETT indispensable à chaque étape du séjour d'un patient en USIC pour ICA
- Echographe dédié
- Le plus tôt possible et le plus souvent possible
- Mesure des pressions de remplissage VG:
Algorithmes +++

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MERCI

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paramètres à retenir

- onde E dépend de la relaxation et de la POG
- TD depend de Pog, Pvg, relaxation VG
- onde A depend contractilité A, PVG, charge OG
- durée onde Apulm-Amitrale (indépendant de l'Evg et de l'age) :
6. Rossvoll O et al. J Am Coll Cardiol 1993 ; 21 : 1687-96.
- mais aussi, TRIV, Flux veineux pulmonaire, Vp, manoeuvres due Valsalva, TD Amitrale, TD onde D pulmonaire, morphologie de la fuite mitrale ou aortique

EVALUATION DES PRESSIONS DE REMPISSAGE VG

TACHYCARDIE SINUSALE = FUSION E et A

E/Ea > 10

TD onde A < 60 ms

Evaluation des PRVG CMH

- $E/E_a > 14$
- Vol OG indexé > 34ml/m²
- Durée Apulm - Amitrale > 30 sec
- Pic IT > 2,8 m/s
- Si > 2 valeurs + = PPRVG augmentées

+/- durée Ap-Am

EVALUATION DES PRESSIONS DE REMPISSAGE VG

AC/FA Onde E exclusive

**TDE < 150 ms
si fonction systolique altérée**

E/Ea > 11

E/ Vp> 2.5

Indépendamment de la fonction systolique

pourquoi mesurer les prvg ?

- en rea d'abord pour évaluer les besoins en remplissage vasculaire chez les pats en défaillance circulatoire
- c'est ce qui conditionne la prise en charge d'un patient dyspnéique voire intubé différenciant l'insuffisance cardiaque d'une autre cause de dyspnée (ep, pneumopathie, sdra, ...) et donc conditionnant la prise en charge thérapeutique que la fonction systolique soit altérée ou non (oap avec fe conservée ça existe mais sera chez un insu cardiaque équilibré ça existe aussi)
- l'echo/doppler permet d'approcher de manière non invasive des mesures hémodynamiques difficile à obtenir de façon invasive
- définition: méthodes invasives déterminer la P diastolique minimale, P moyenne, la PTDVG pré A et PTDVG, toutes ces valeurs n'évoluent pas forcément parallèles; le plus souvent les corrélations sont faites avec Pog moyenne
- pronostic: FEvg basse (CMD ou post IDM) et profil restrictif ; bas

9. Rihal CS et al. Circulation 1994 ; 90 : 2772-9. Rechercher l'
10. Nijland F et al. J Am Coll Cardiol 1997 ; 30 : 1618-24.

INsuf cardiaque droite aigue rôle de l'echo

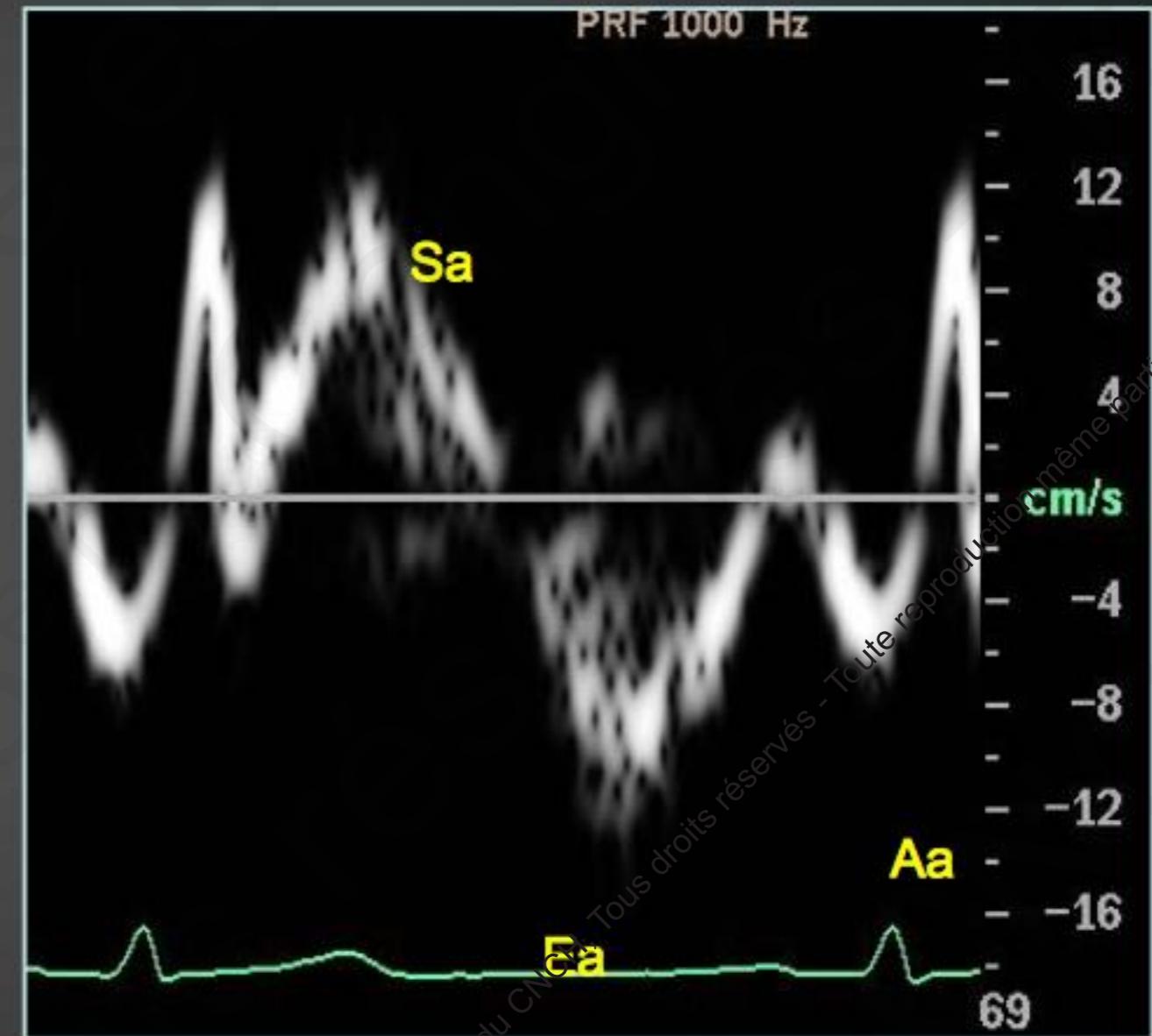
Fonction ventriculaire droite

- morphologie
- cinétiqe pariétale
- fonction systolique
- diastole
- conditions de charge

Fonction VD: *DTI anneau tricuspidien*



Sa > 15 cm/s



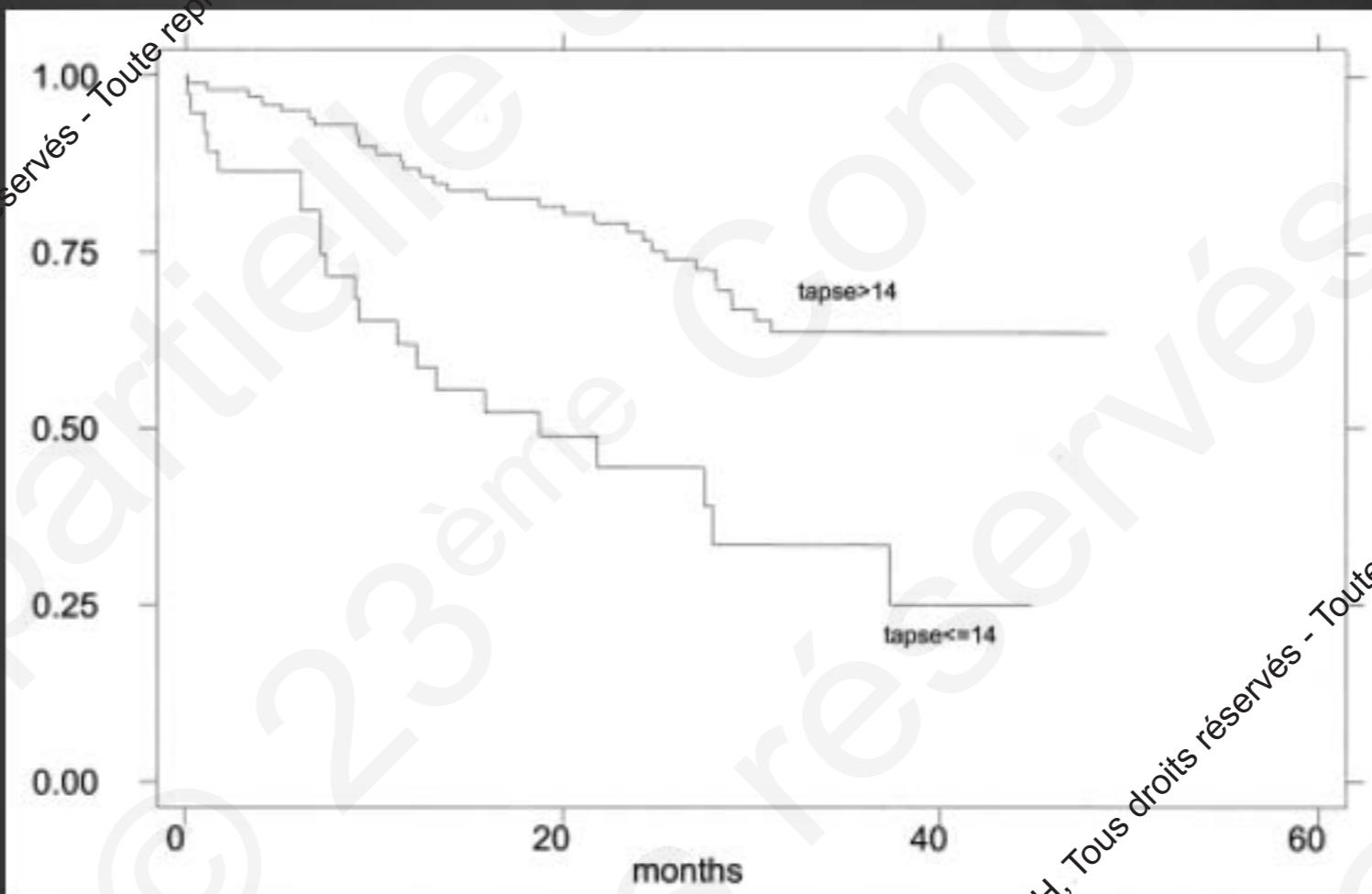
Fonction VD: *TM anneau tricuspidien*



normale ≥ 16mm

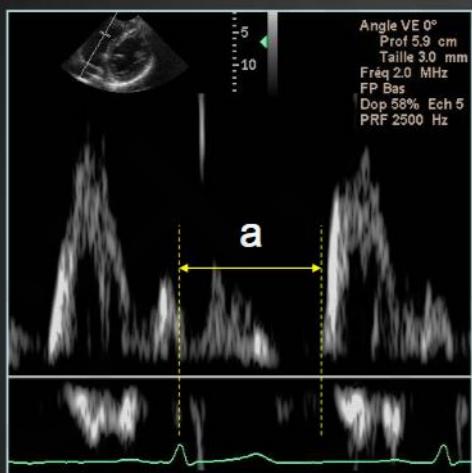
dysfonction VD: < 12mm (r = 0,92 FE isotopique)

Ventricule droit – TAPSE: pronostic

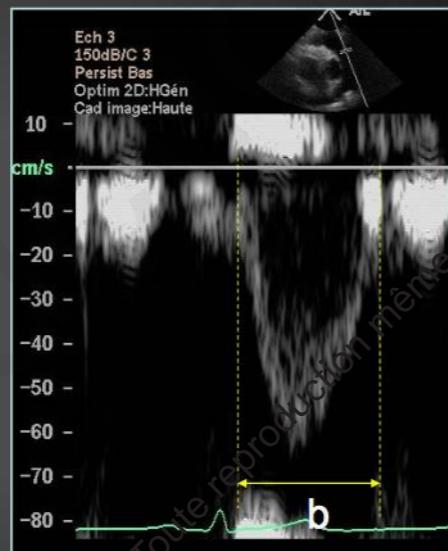


Ghio AJC 2000

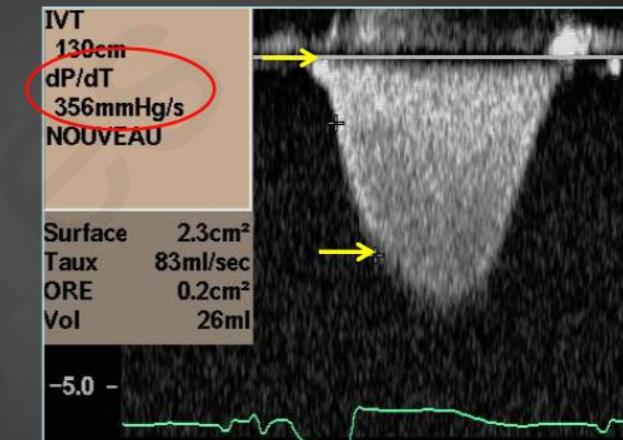
Ventricule droit: index de Tei



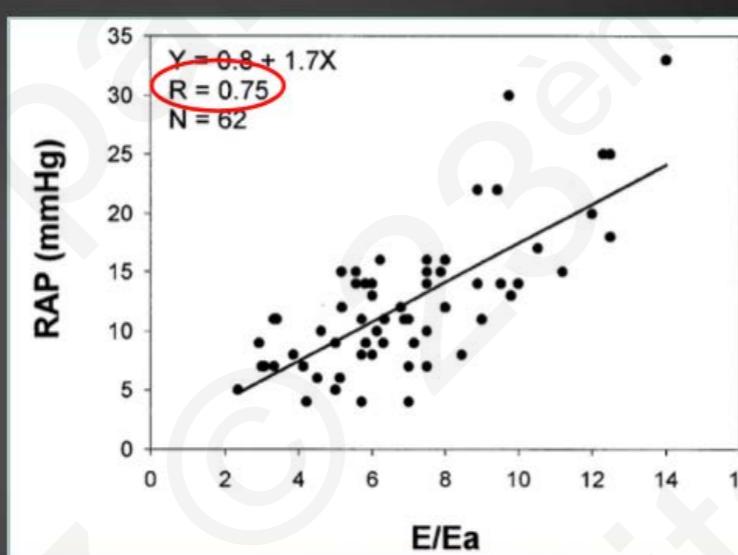
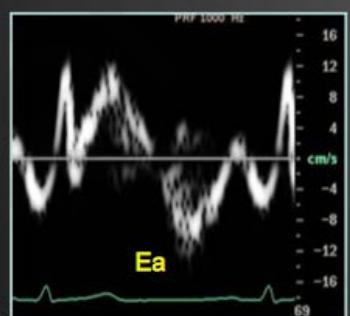
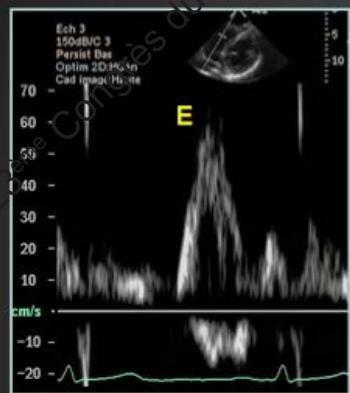
normal: 0.28 ± 0.04



Fonction VD: $IT - dP/dT$



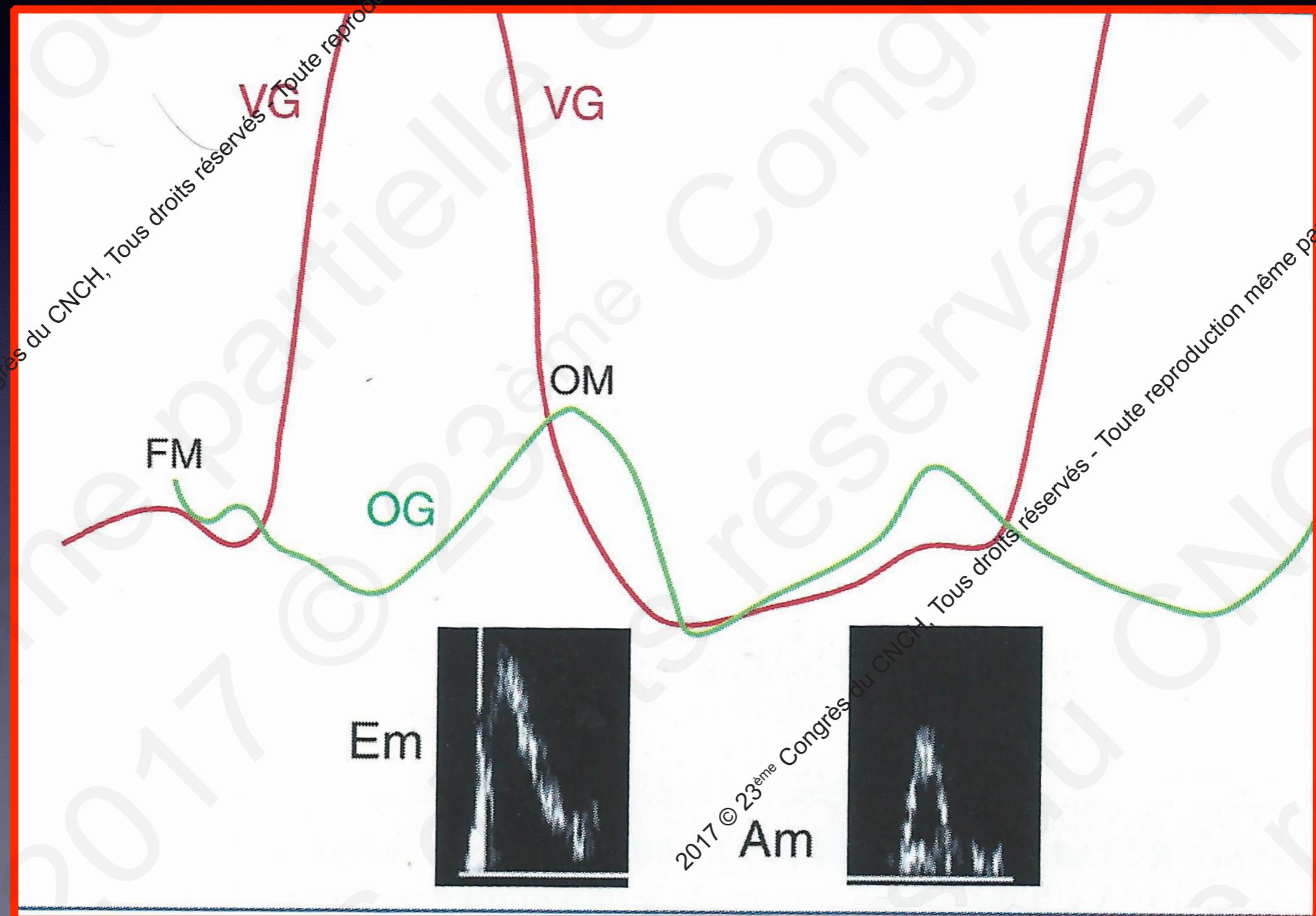
Pression OD: *doppler tricuspidien / doppler tissulaire anneau*



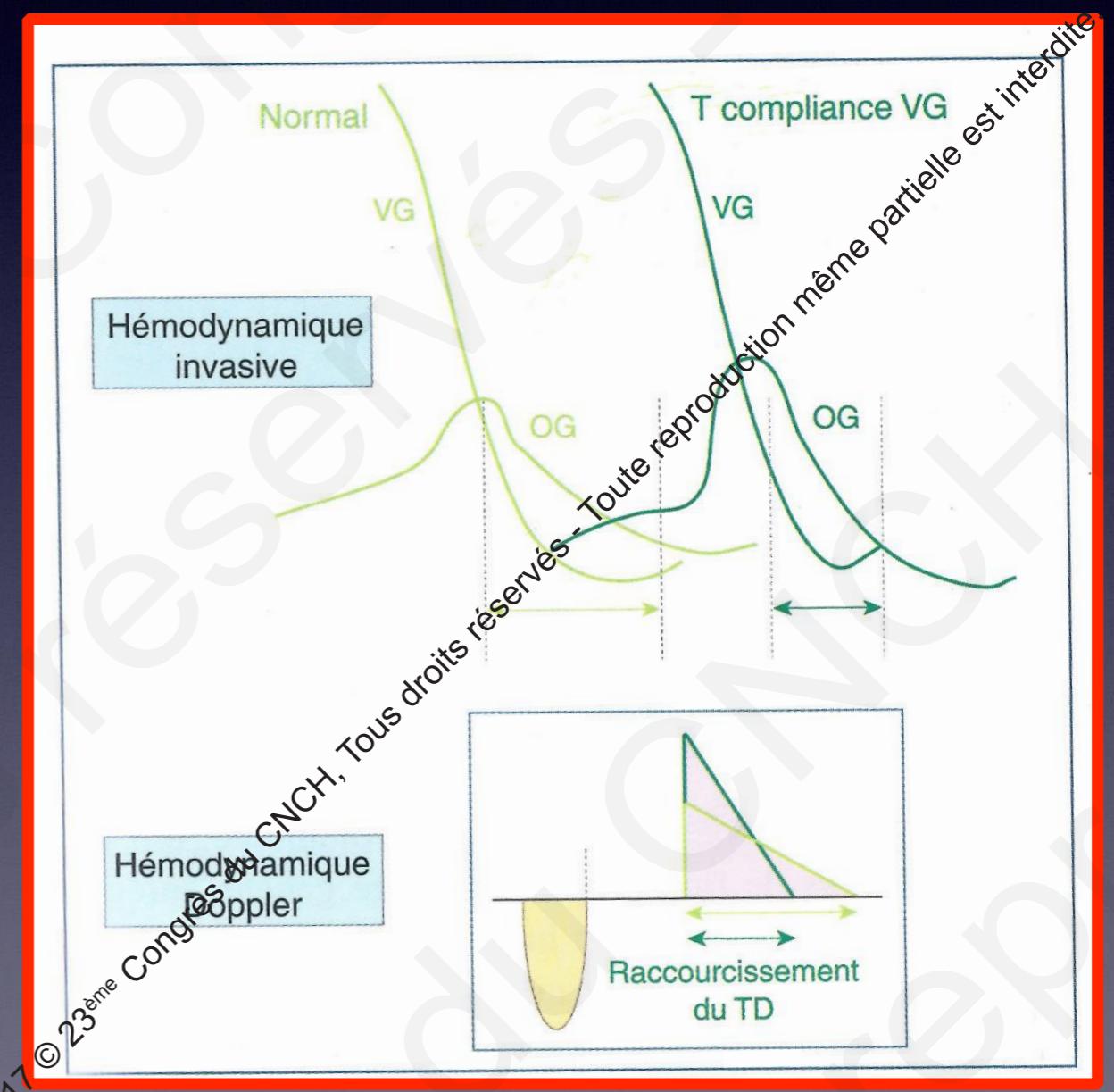
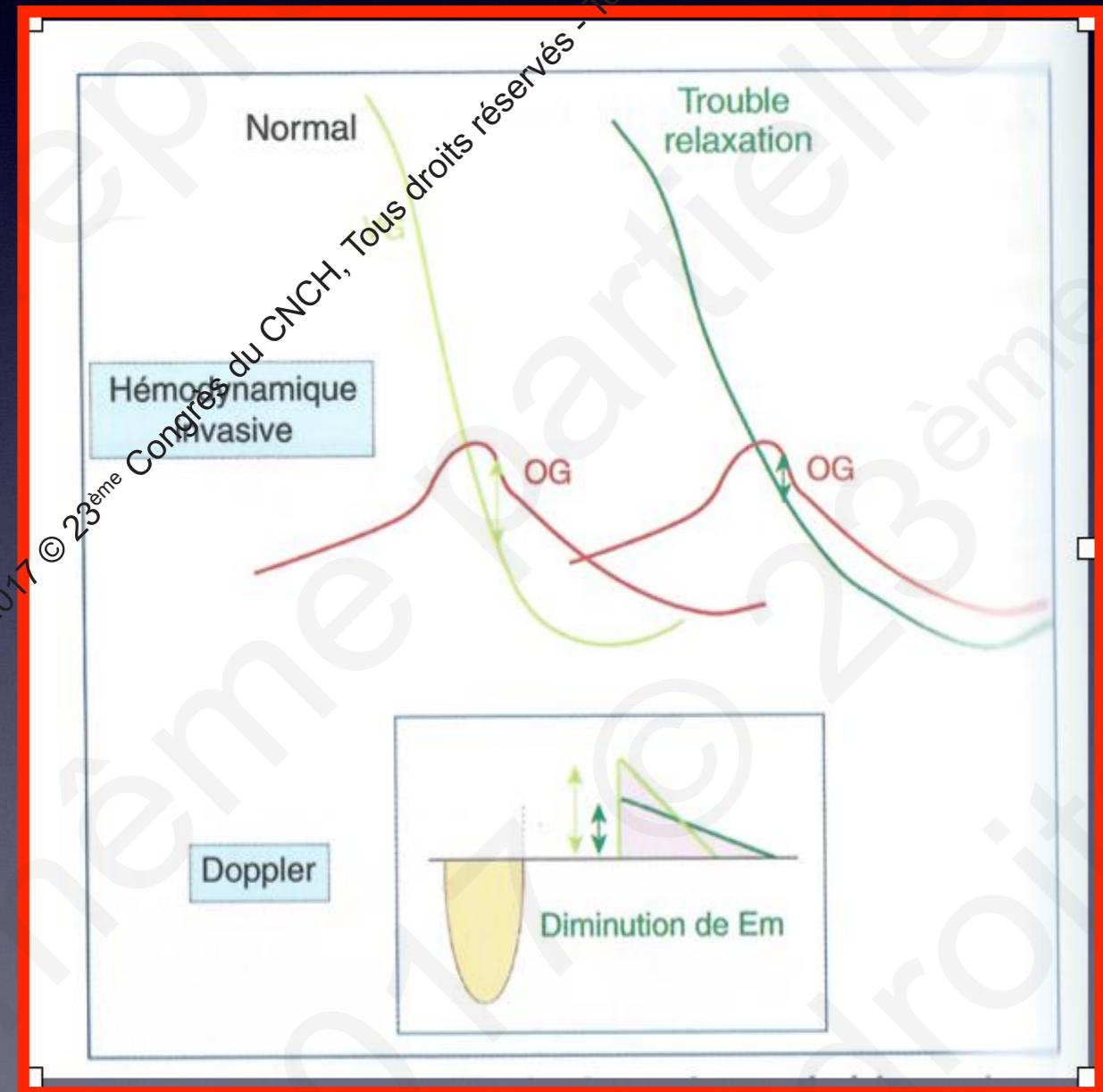
$E/Ea > 6 \rightarrow P\ OD \geq 10\ mm\ Hg$ (Se = 79% - Sp = 73%)

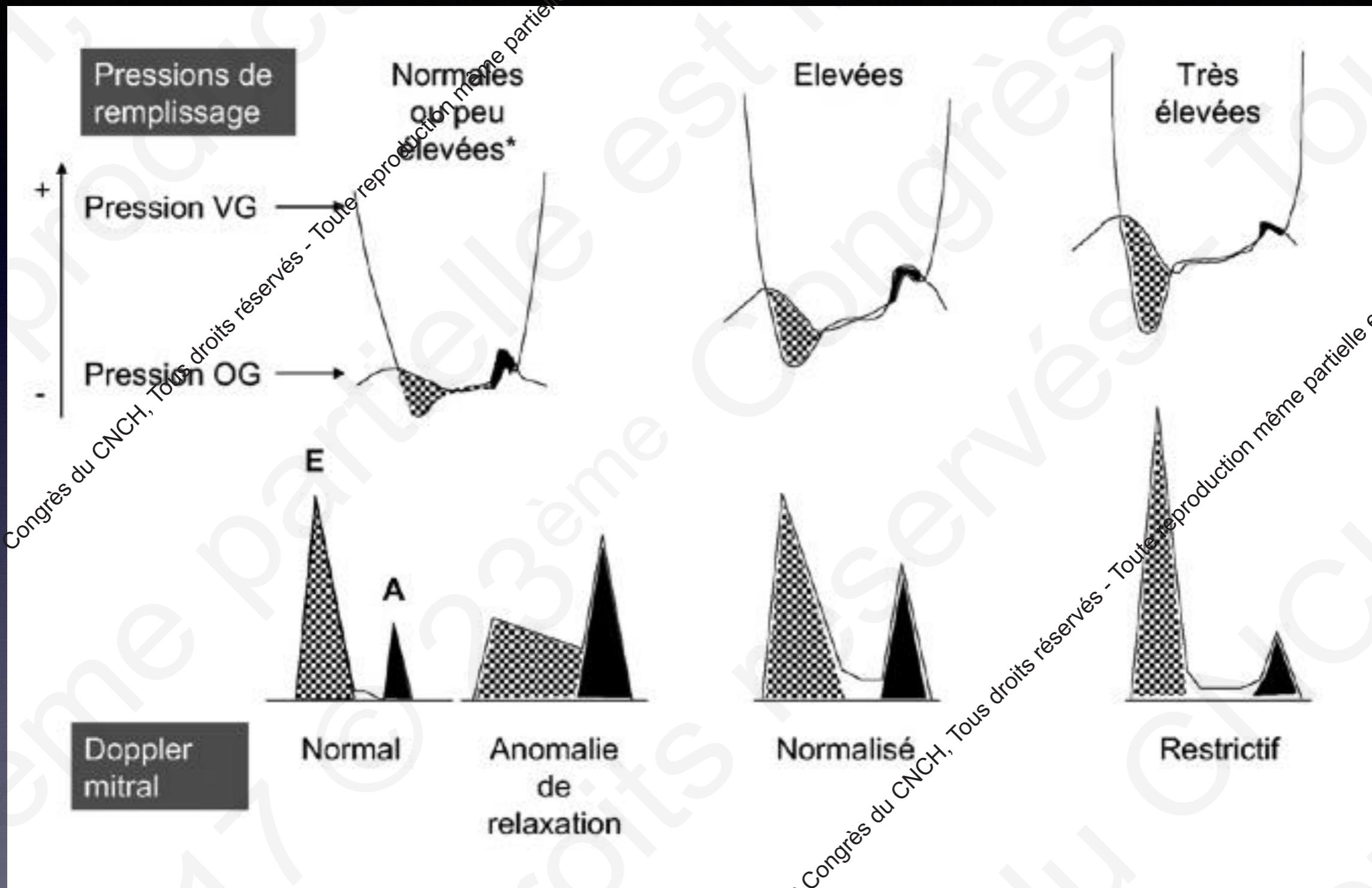
Nagueh Am J Cardiol 1999

Flux transmitral normal



Onde E et TD





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