

# SIMA 23

Σι-μα Ιητηρ(ιατρός) Μινωϊκή Κρήτη

1ο ΠΑΝΕΛΛΗΝΙΟ  
ΠΟΛΥΘΕΜΑΤΙΚΟ ΣΥΝΕΔΡΙΟ  
ΙΑΤΡΙΚΟΥ ΣΥΛΛΟΓΟΥ ΗΡΑΚΛΕΙΟΥ

ΣΥΝΔΙΟΡΓΑΝΩΣΗ:



ΠΕΡΙΦΕΡΕΙΑ ΚΡΗΤΗΣ  
REGION OF CRETE



## νόσο και αιφνίδιος καρδιακός θάνατος. Αλγόριθμοι αντιμετώπισης

Σ. Ζερβάκης

MD, Ειδικευόμενος Καρδιολογίας Πα.Γ.Ν.Ηρακλείου

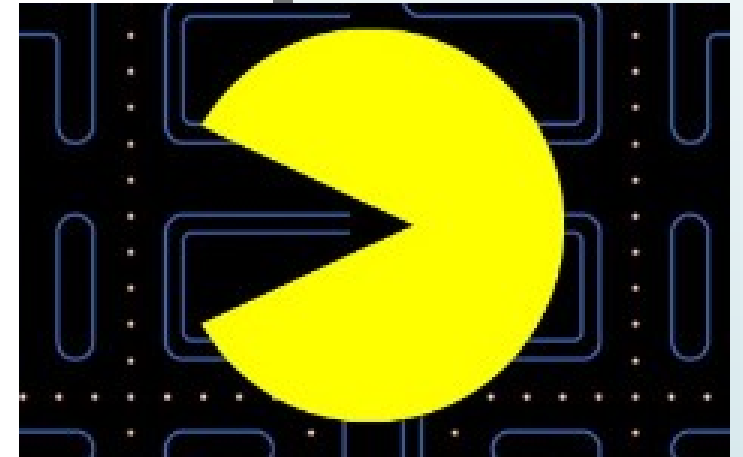
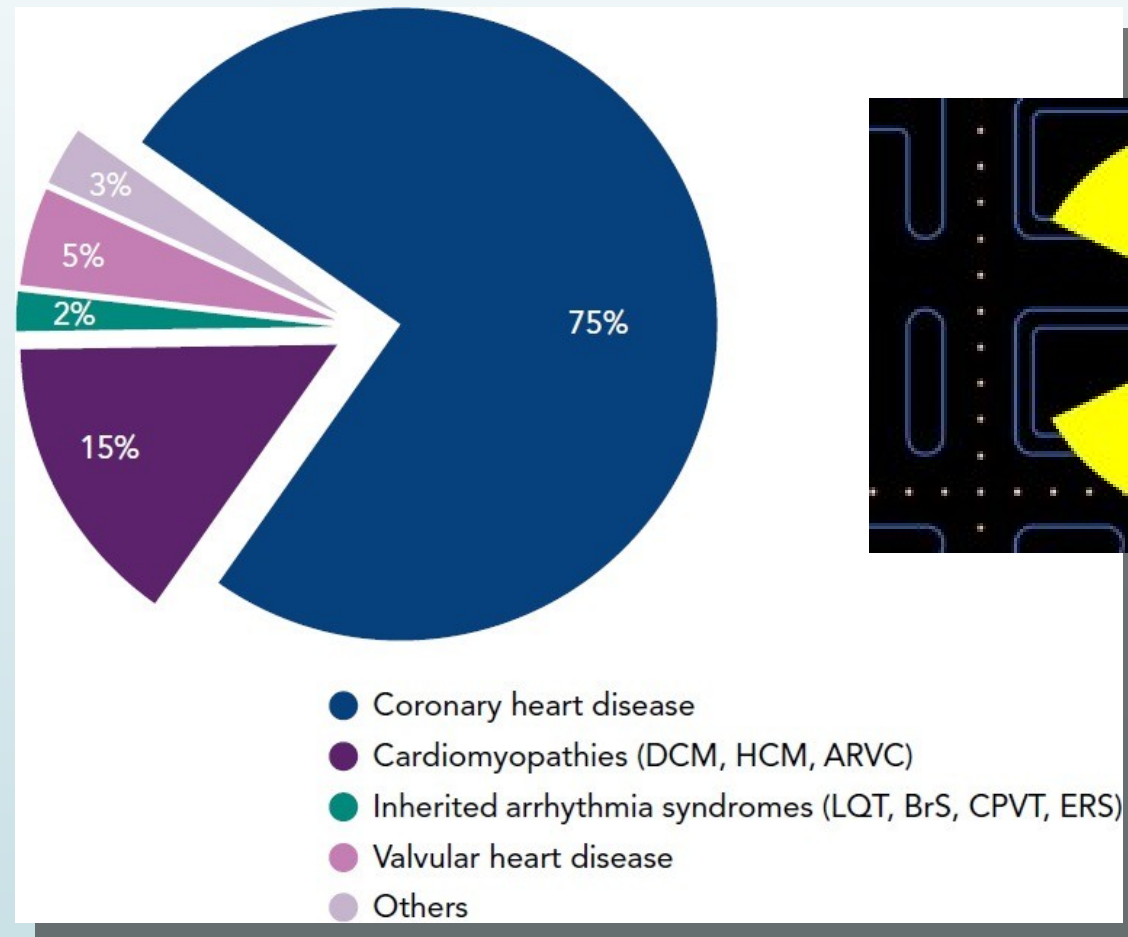


Αιφνίδιος καρδιακός θάνατος και...

... στεφανιαία νόσος

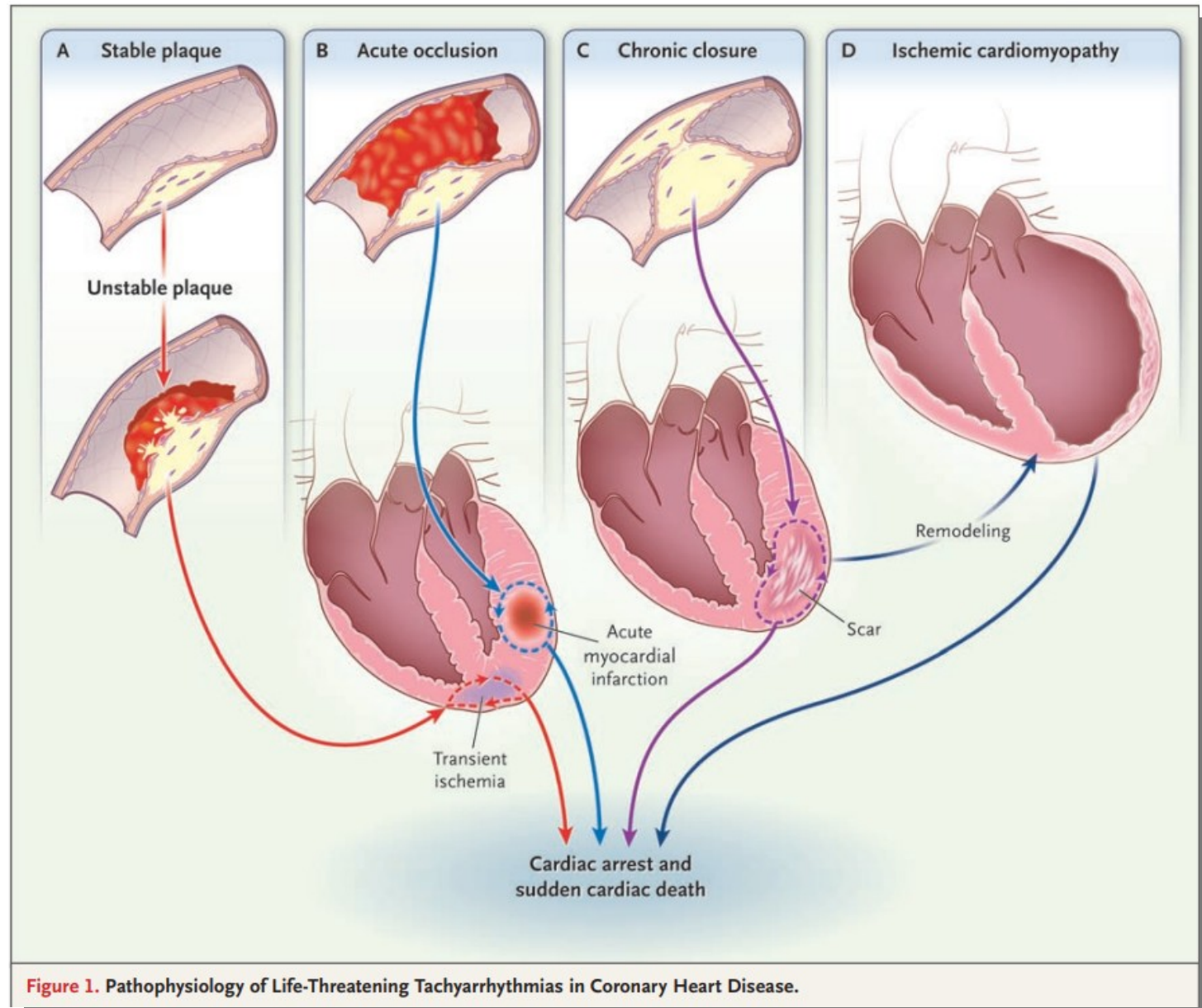
Ευθύνεται συνολικά για τη **συντριπτική πλεινότητα** των αιφνιδίων θανάτων στον Δυτικό κόσμο

Στεφανιαία νόσος



# Στενή σχέση μεταξύ

- στεφανιαίας νόσου
- και αιφνιδίου θανάτου



**Table 1.** Primary modes of cardiac arrest/SCD and mechanisms in selected disorders

<b>Disorder</b>	<b>Primary mode of cardiac arrest/SCD</b>	<b>Mechanism</b>
Brugada syndrome	VT/VF	Reflection (phase 2 reentry)
CAD (Acute ischemia)	VT/VF	Multiple (reentry, automaticity, triggered activity)
CAD (Prior MI)	VT	Scar-mediated reentry
CPVT	VT/VF	Delayed after depolarizations
Dilated cardiomyopathy	VT	Scar-mediated reentry Bundle branch reentry
Hypertrophic cardiomyopathy	VT/VF	Multiple
Long QT syndrome	Torsades de pointes VT	Early after depolarizations
WPW syndrome	VF	Rapid conduction to the ventricles down an accessory pathway resulting in VF

*CAD*, coronary artery disease; *MI*, myocardial infarction; *VT*, ventricular tachycardia; *VF*, ventricular fibrillation; *CPVT*, catecholaminergic polymorphic ventricular tachycardia

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**Table 1. Effects of Metabolic Changes on Cardiac Ion Channel/Transporter Function and Arrhythmogenicity**

Channel/Transporter Effects	Metabolic Changes	Effects on Electric/Ionic Homeostasis	Proarrhythmic Mechanism
Na <sup>+</sup> /K <sup>+</sup> ATPase ↓	Ischemia/hypoxia	Na <sup>+</sup> overload	Ca <sup>2+</sup> overload and DAD
Cx hemichannel ↑	Ischemia	Na <sup>+</sup> overload	Ca <sup>2+</sup> overload and DAD
Peak I <sub>Na</sub> ↓	Ischemia/heart failure	↓ Na <sup>+</sup> influx	Slow conduction
Late I <sub>Na</sub> ↑	Ischemia/hypoxia acidosis, ↑ LPC AMPK mutation	↑ Na <sup>+</sup> influx, prolonged APD	EAD
Kv ↓	Diabetes mellitus, heart failure	↓ K <sup>+</sup> influx, prolonged APD	EAD
Kv ↑	Insulin treatment in diabetic heart, PI3Kα activation, exercise training	↑ K <sup>+</sup> channel expression	Protective
I <sub>KATP</sub> ↑	Ischemia	↑ K <sup>+</sup> influx, shortened APD	Current sink, slow conduction
RyR2	↓ during ischemia, ↑ on reperfusion	SR Ca <sup>2+</sup> load ↑, spontaneous Ca <sup>2+</sup> waves ↑	Ca <sup>2+</sup> transient/action potential alternans

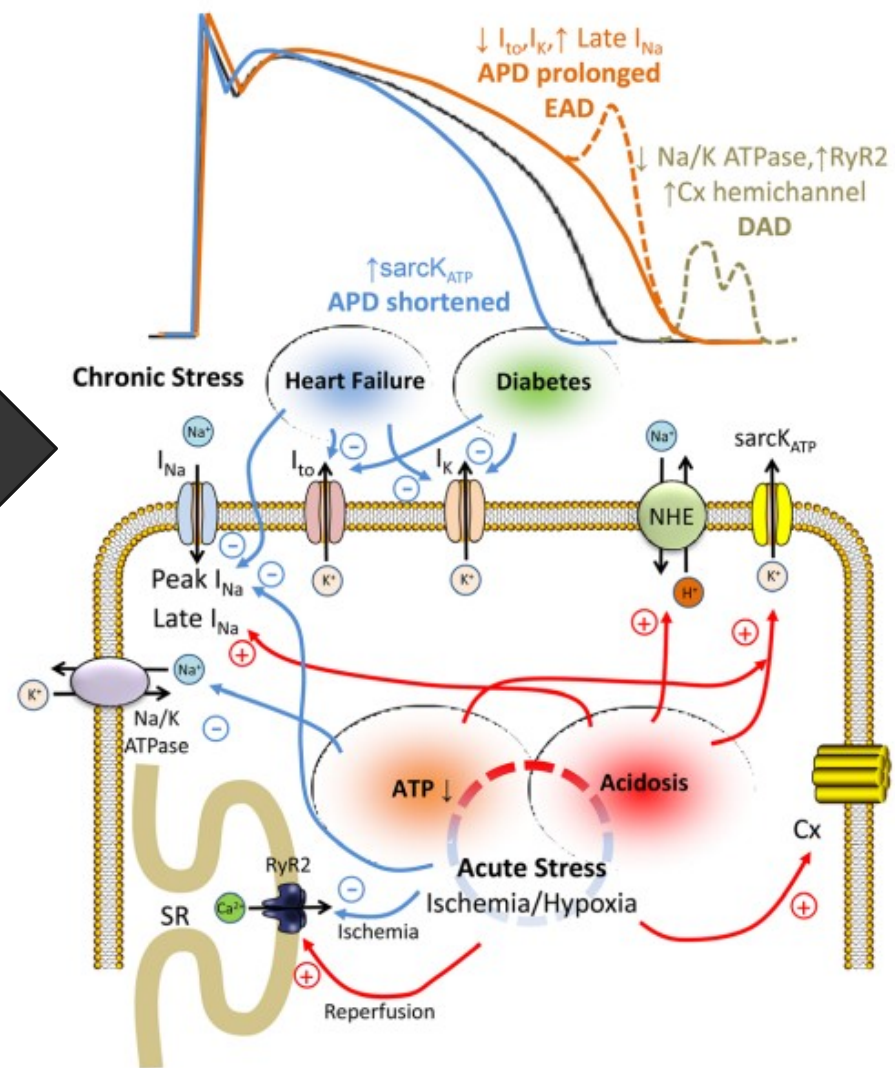
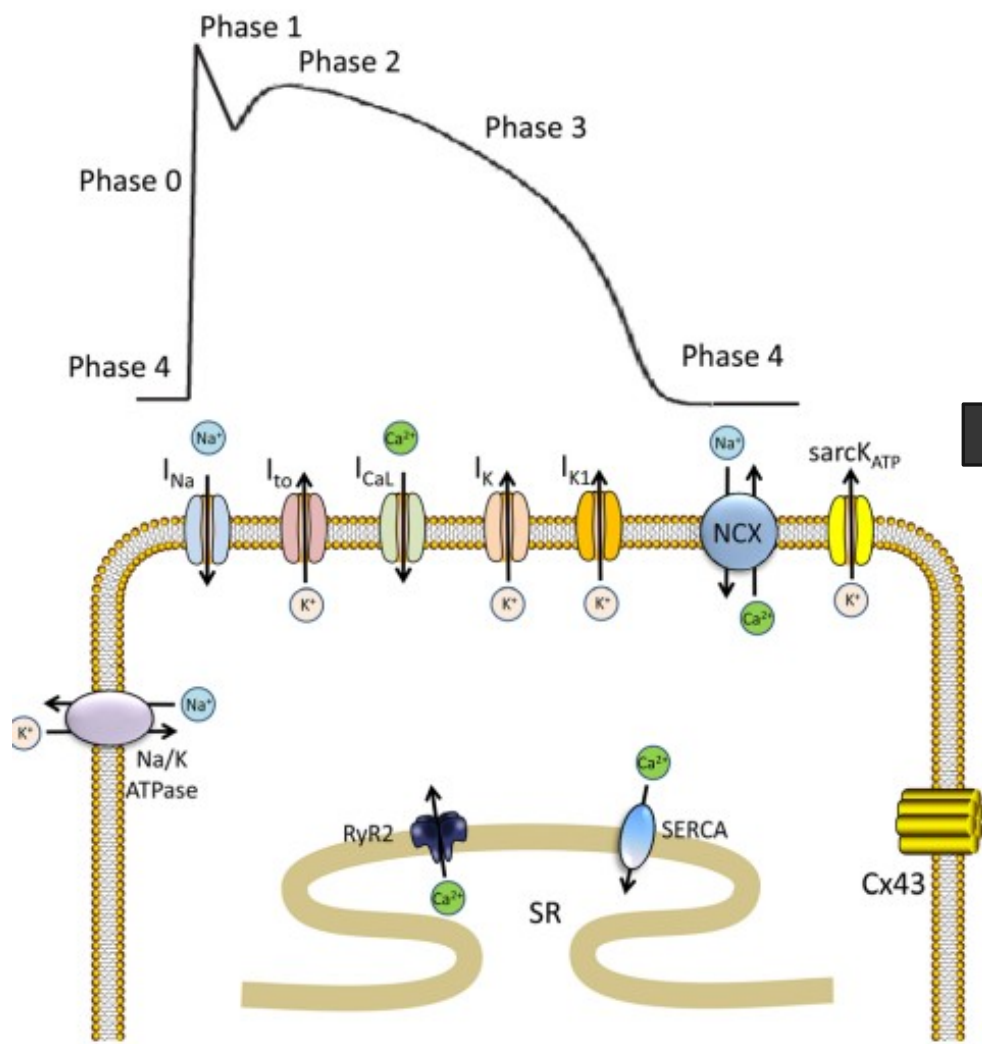
APD, action potential duration; Cx, connexin; DAD, delayed afterdepolarization; EAD, early afterdepolarization; I<sub>KATP</sub>: ATP-sensitive K<sup>+</sup> current; Kv, voltage-gated K<sup>+</sup> current; Late I<sub>Na</sub>, late Na<sup>+</sup> current; LPC, lysophosphatidylcholine; Peak I<sub>Na</sub>, peak Na<sup>+</sup> current; RyR2, ryanodine receptor 2; and SR, sarcoplasmic reticulum.

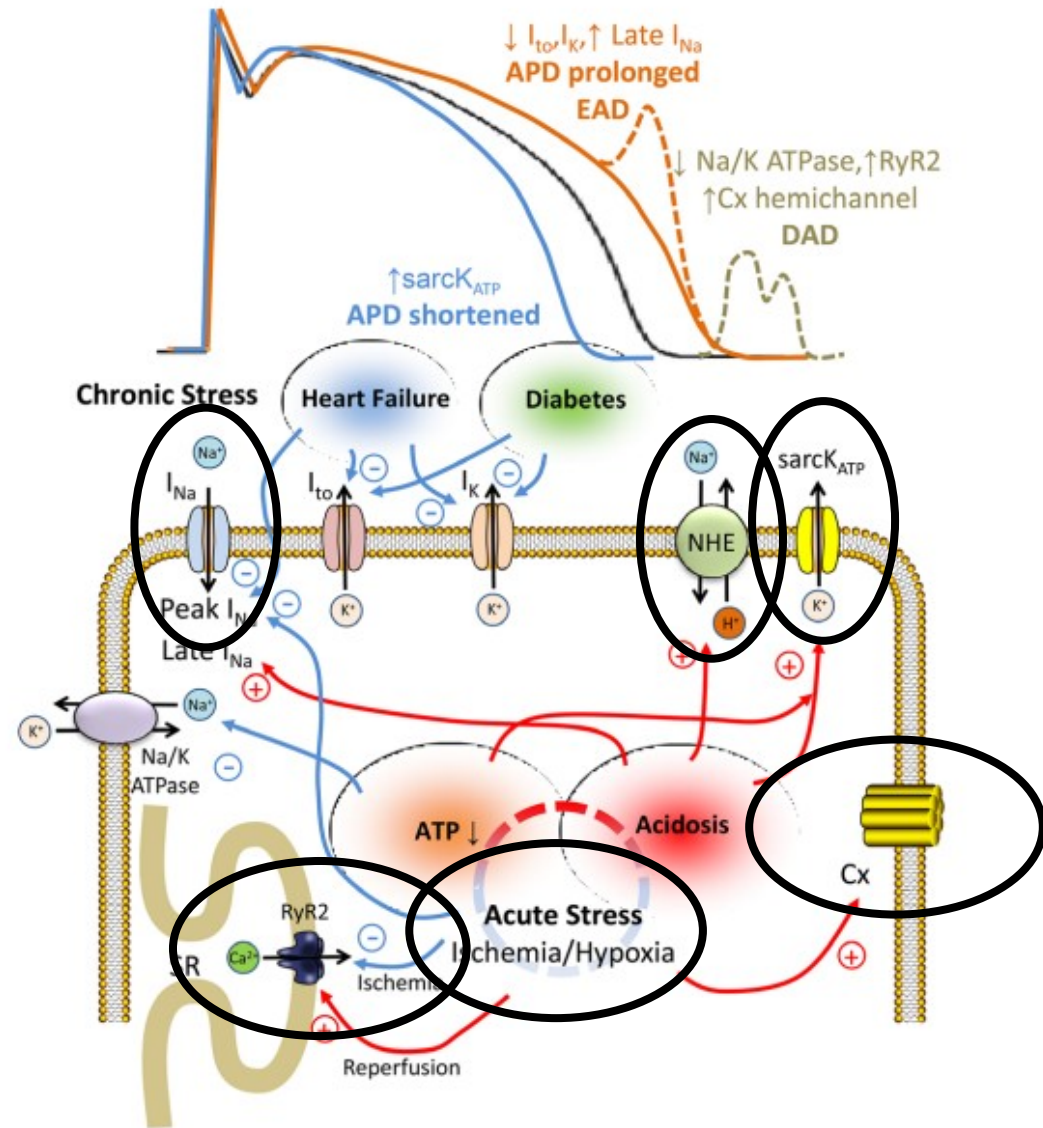
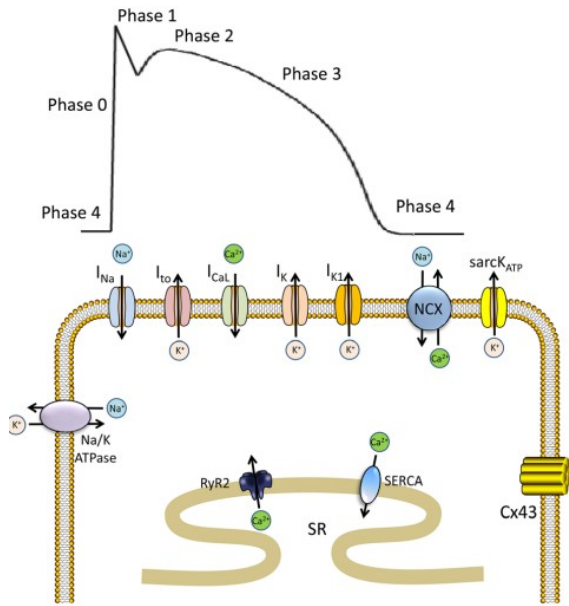


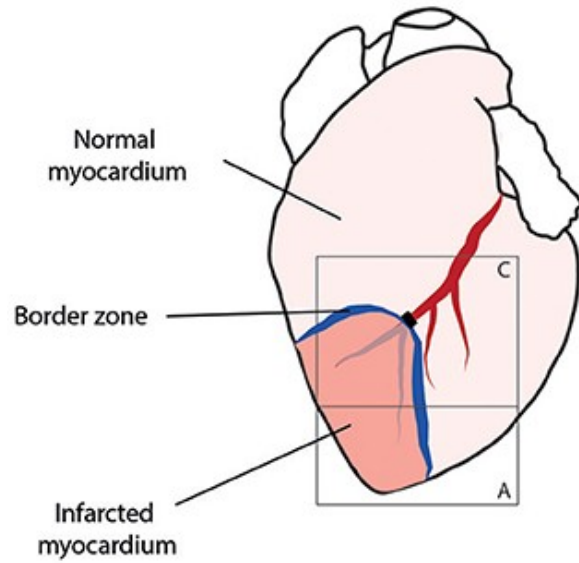
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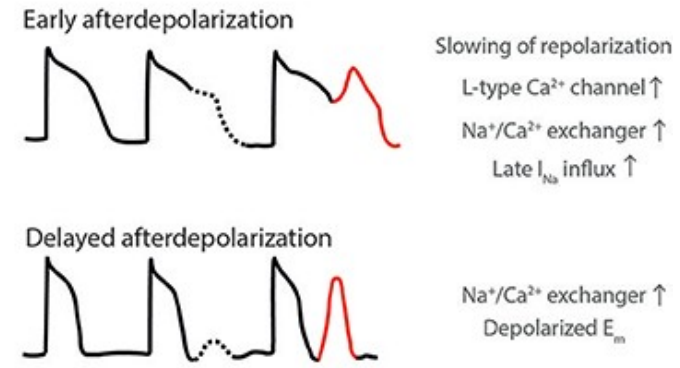
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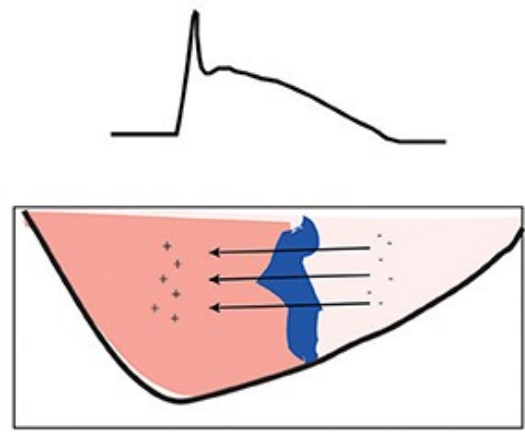




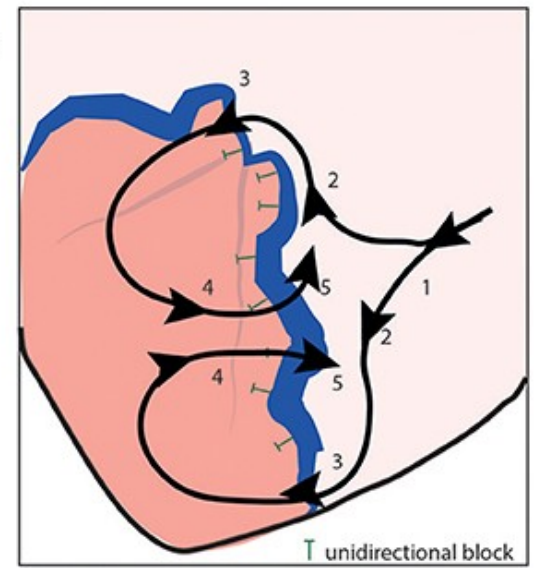
**B** Triggered activity



**A** Automaticity (Injury current)



**C** Reentry



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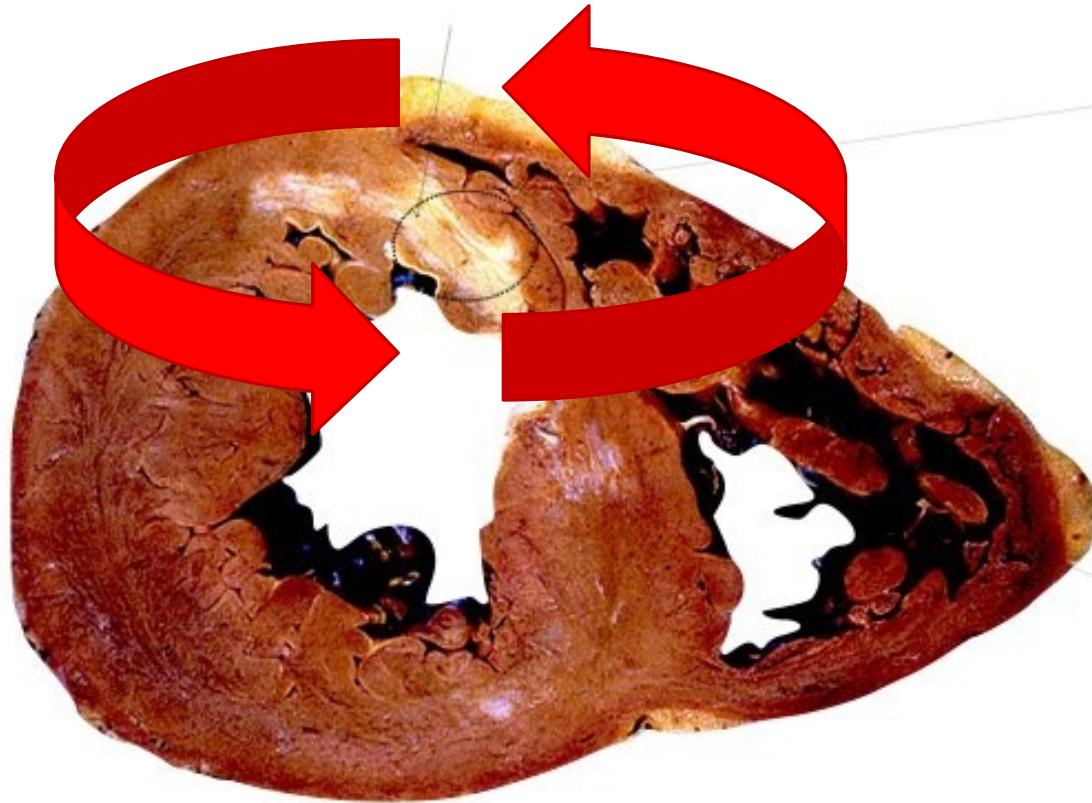
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Established Myocardial Infarction leading to a scar

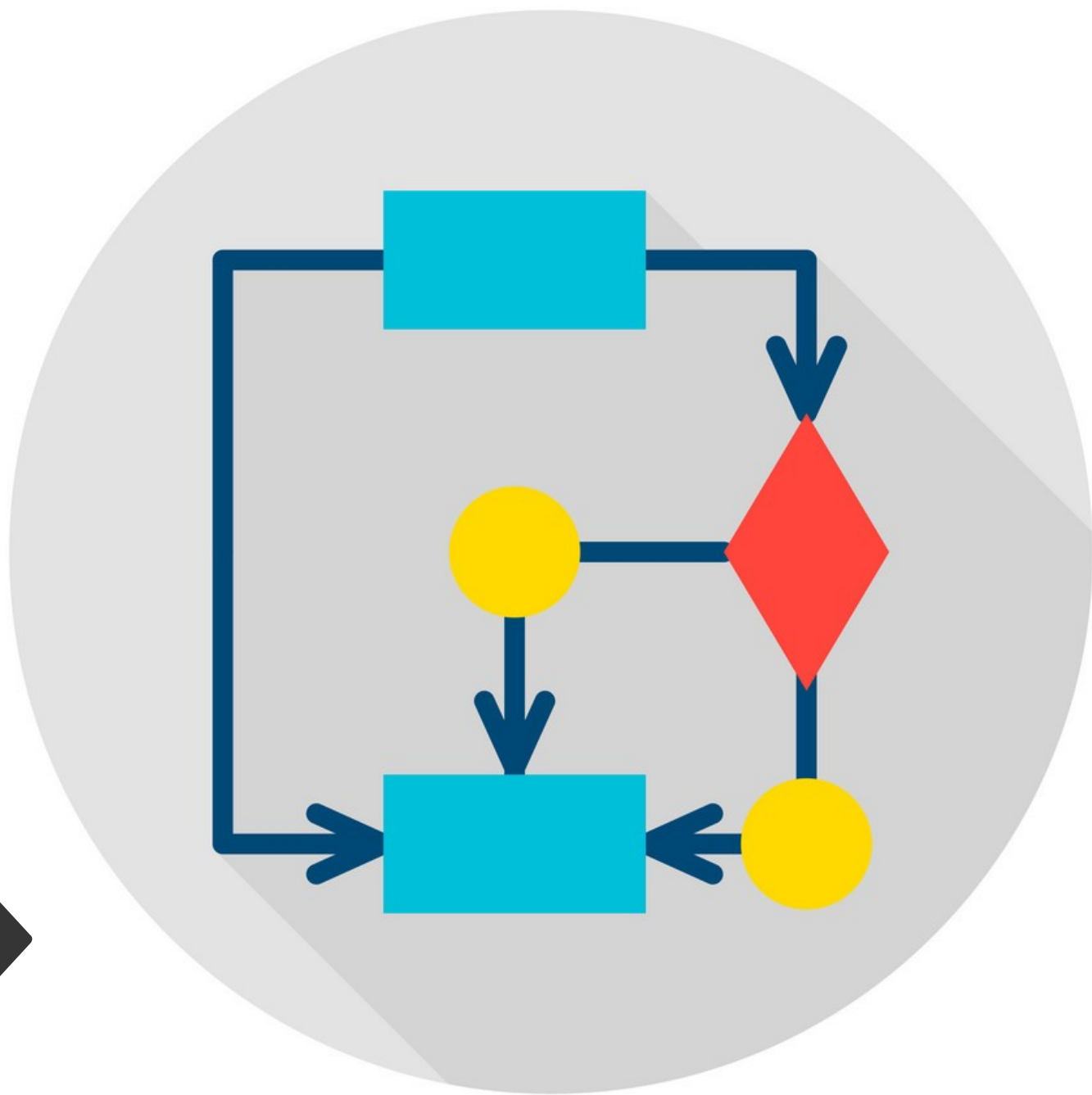


Myocardial Infarction



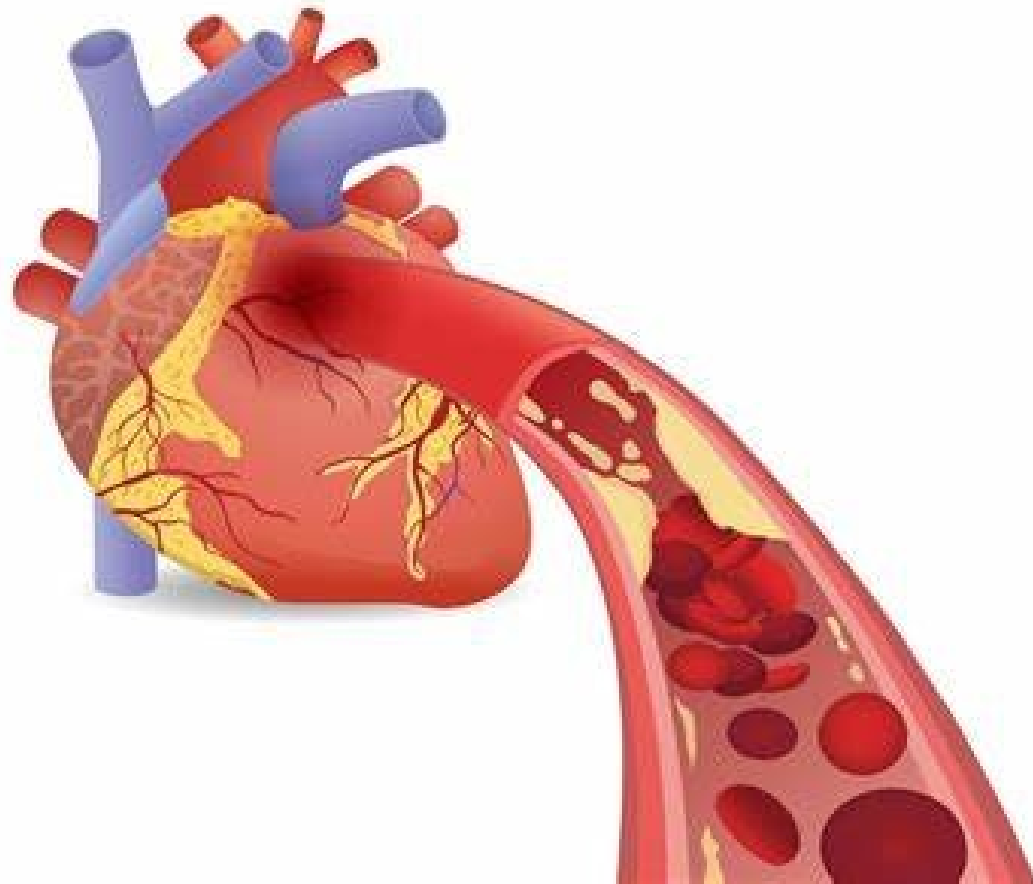
Schematic up of scar tissue

# Αλγόριθμοι αντιμετώπισης



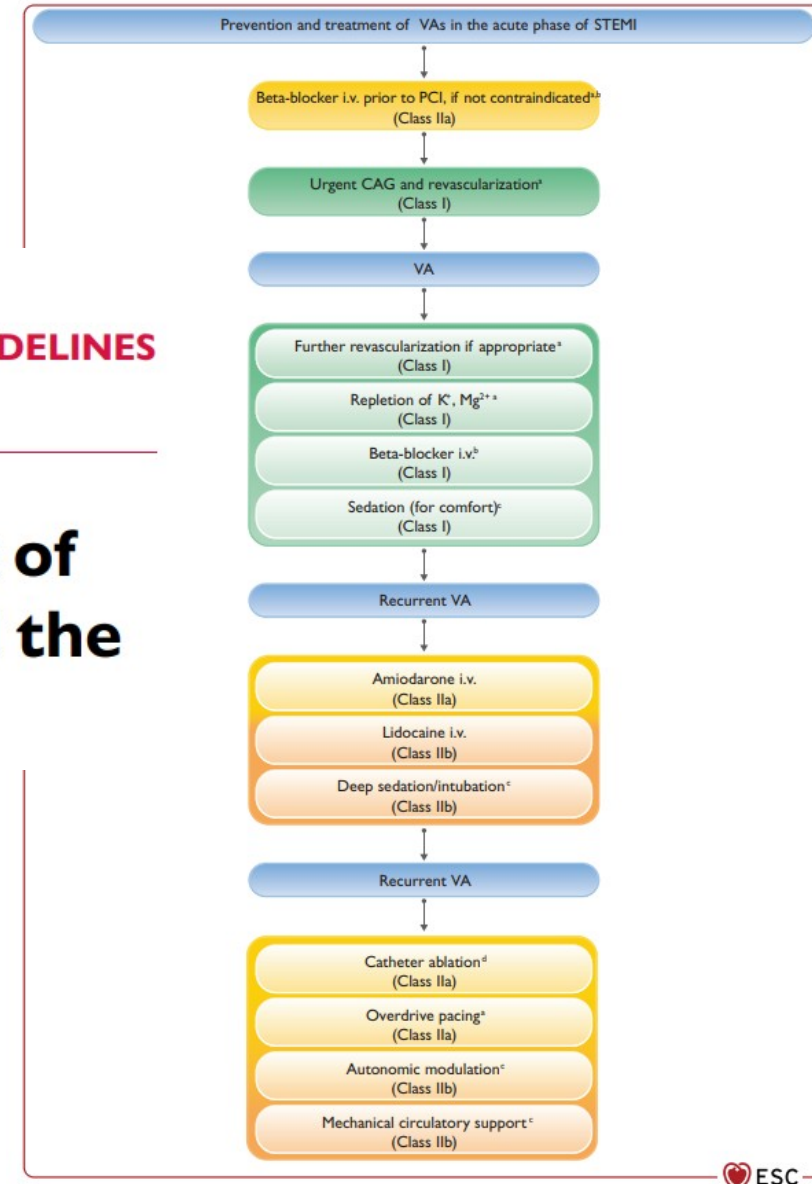


# Οξεία ισχαιμία



**4–12%** των STEMI → κοιλιακή  
αρρυθμιογέννεση το πρώτο 48ωρο  
(κυρίως το πρώτο 24ωρο)

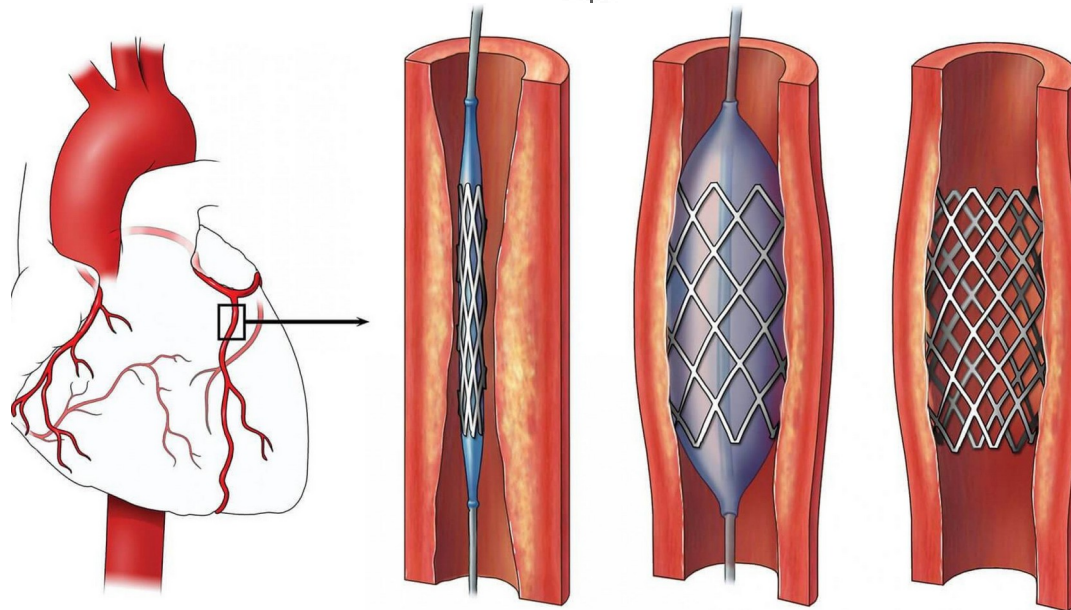
# 2022 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death



Prevention and treatment of VAs in the acute phase of STEMI

Beta-blocker i.v. prior to PCI, if not contraindicated<sup>1,2</sup>  
(Class IIa)

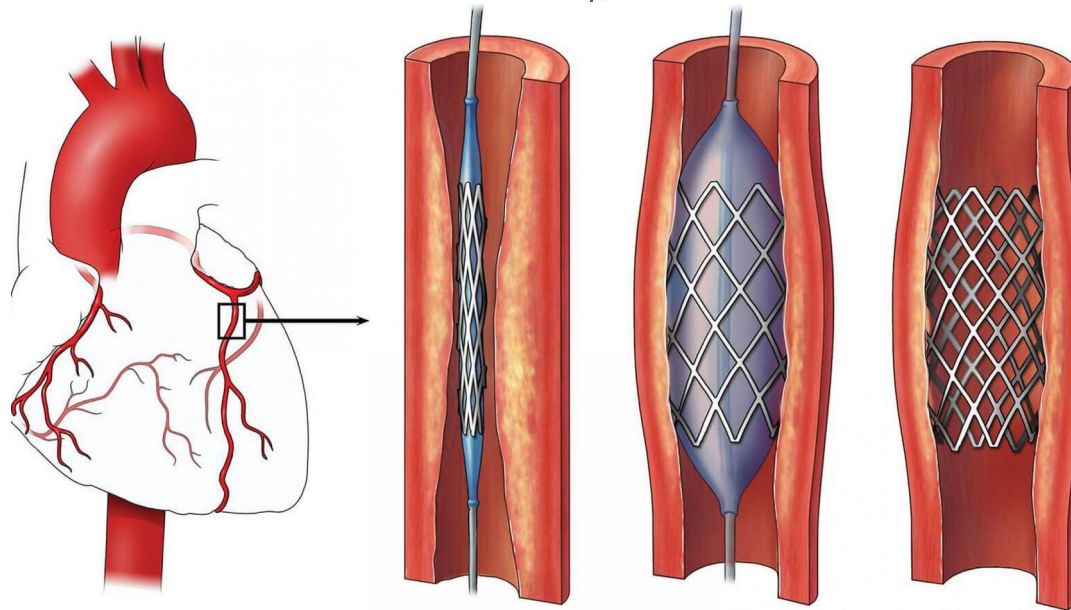
Urgent CAG and revascularization<sup>3</sup>  
(Class I)

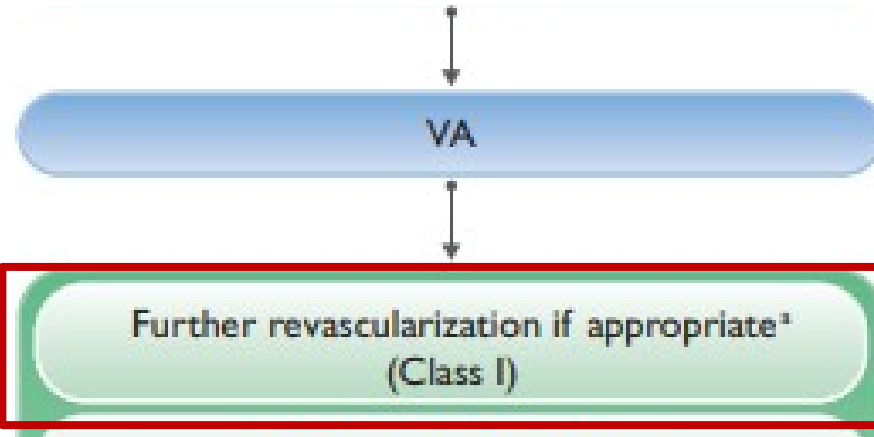


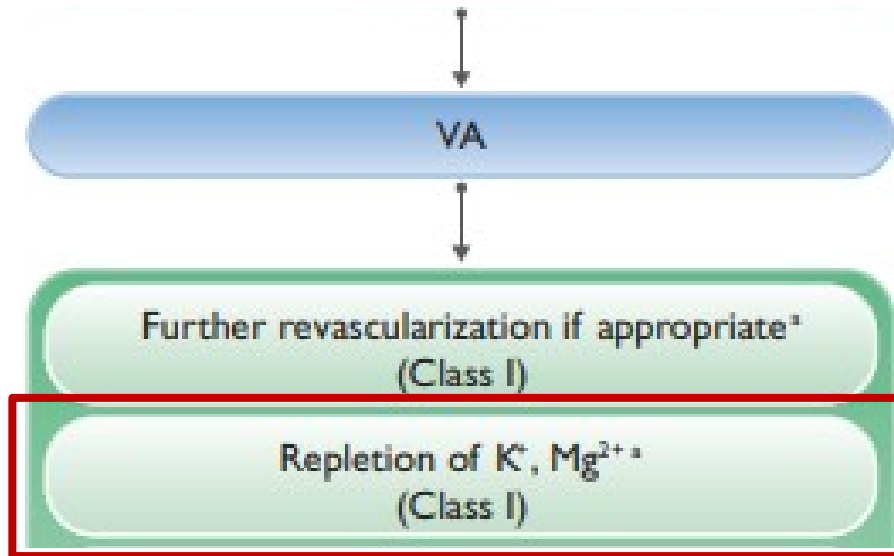
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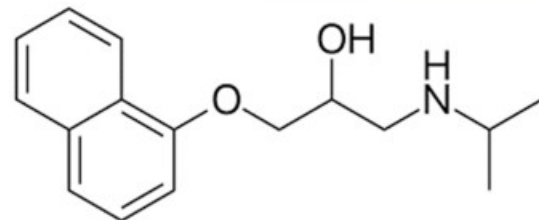
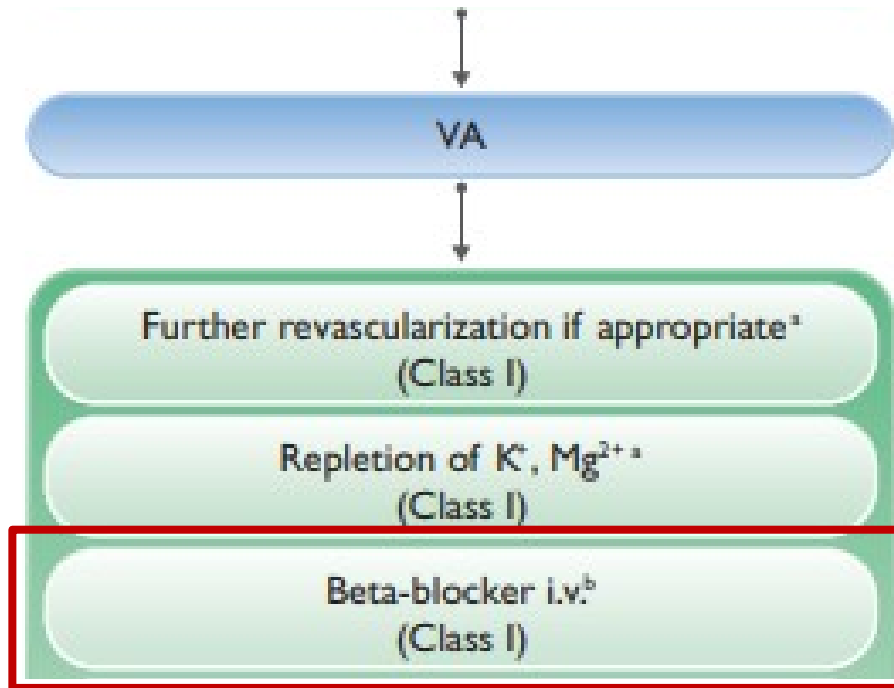
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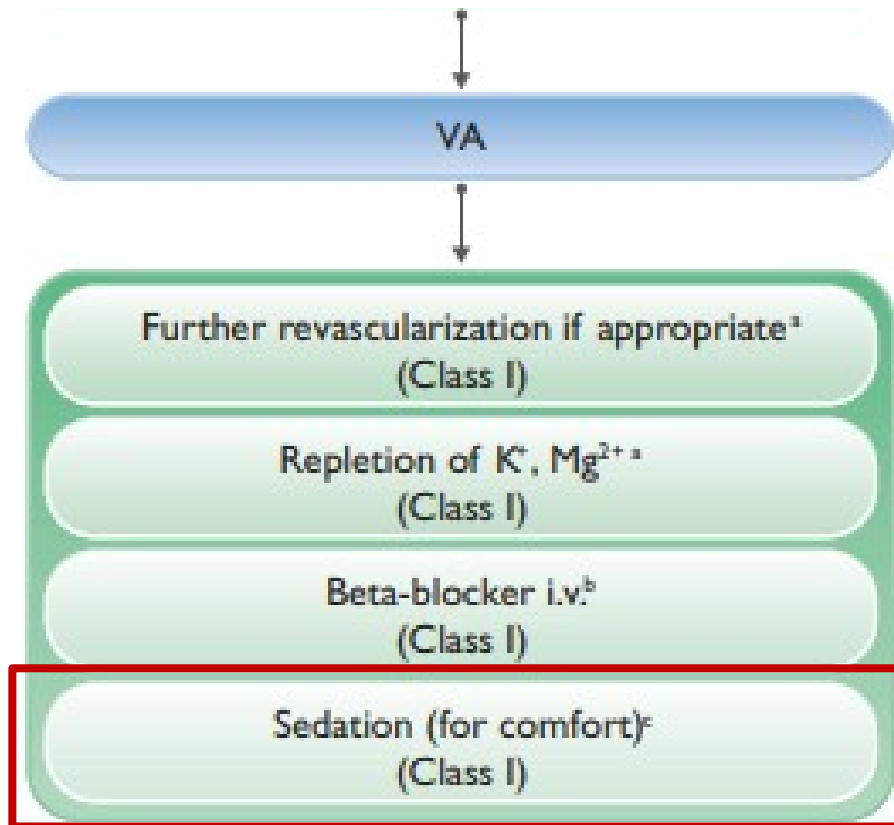
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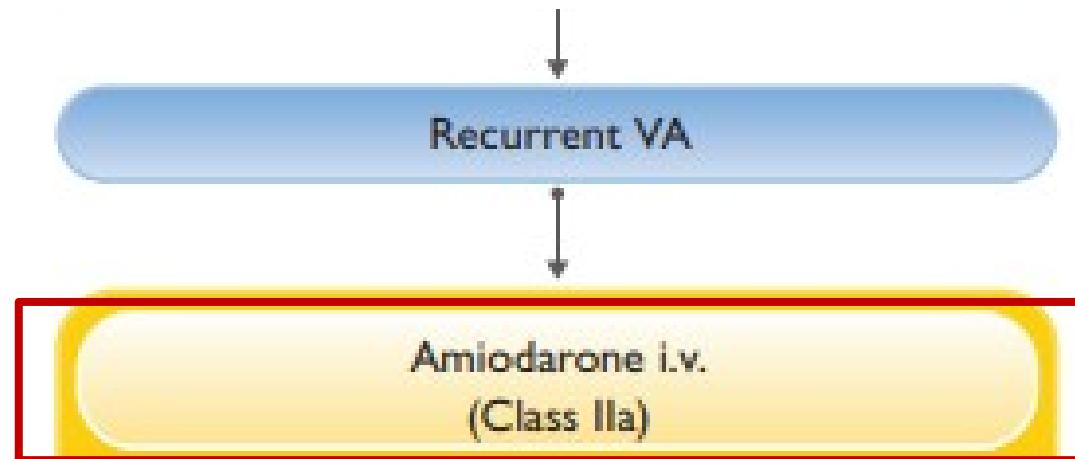






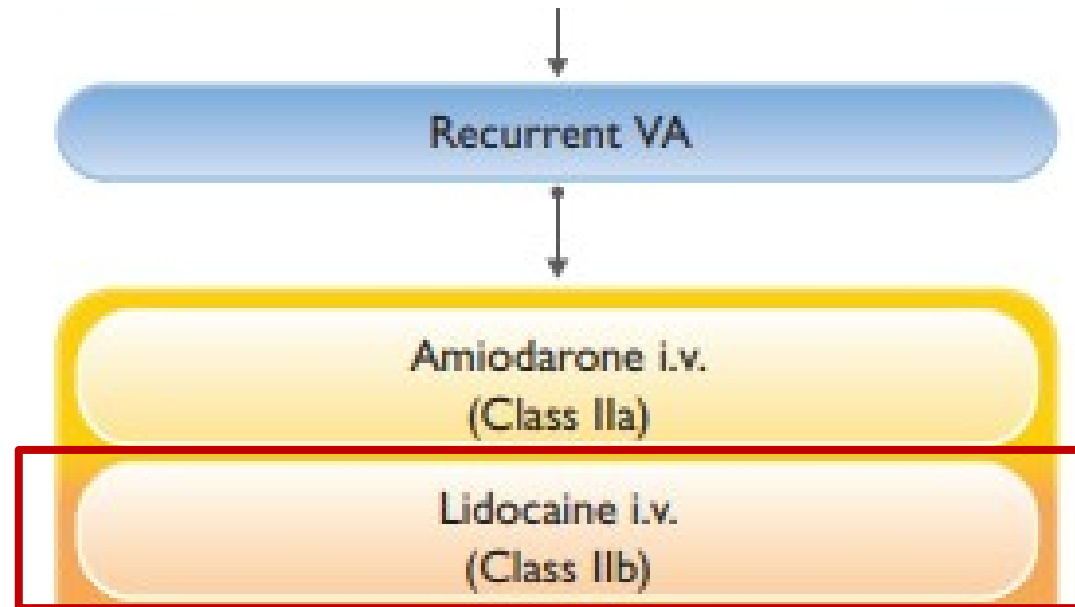






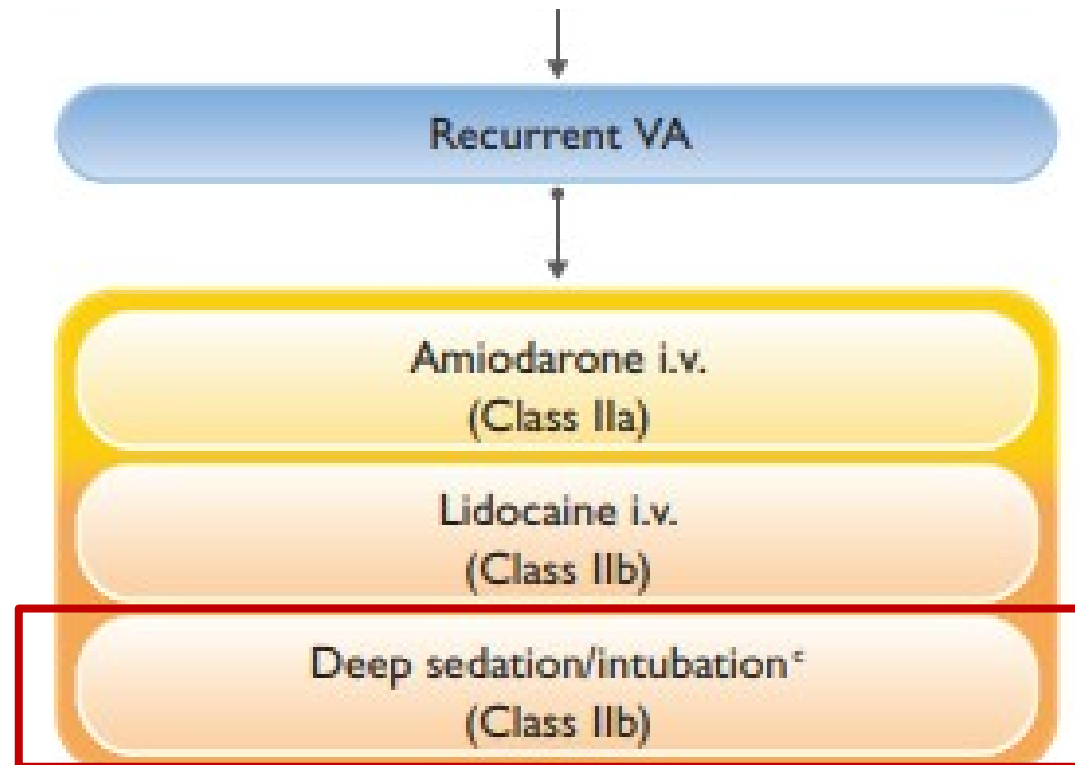
**Table 8** Anti-arrhythmic drugs (acute and chronic treatment)

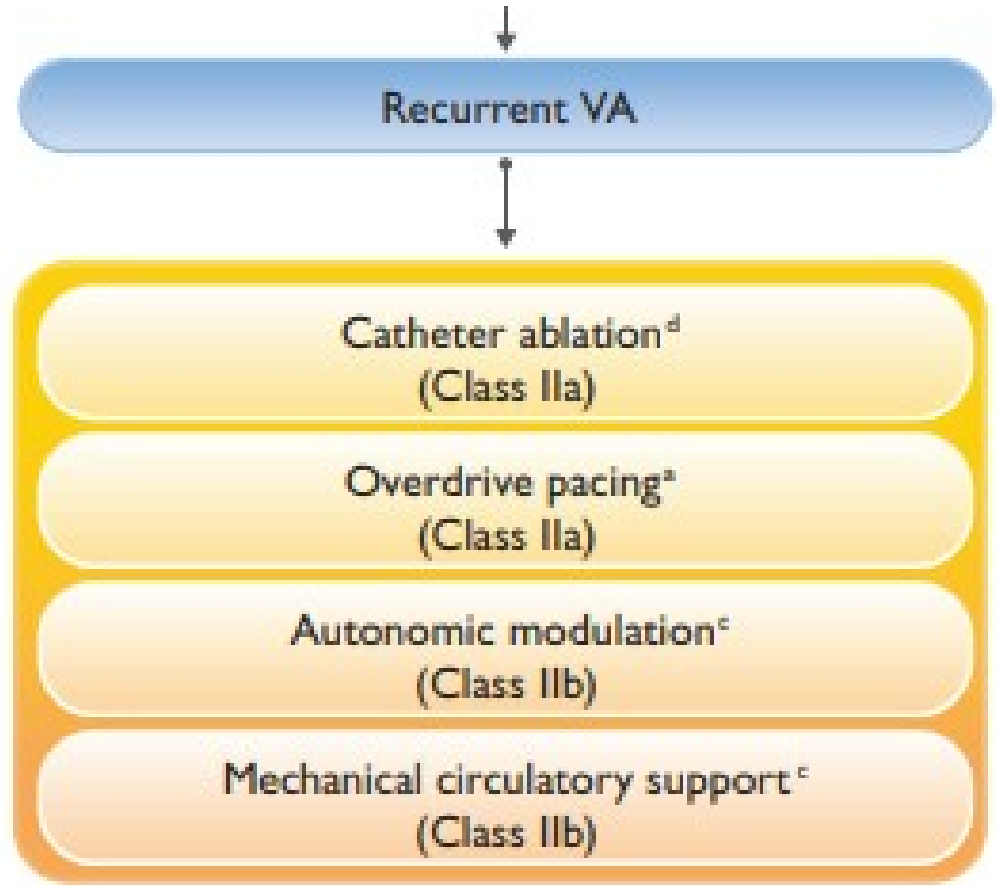
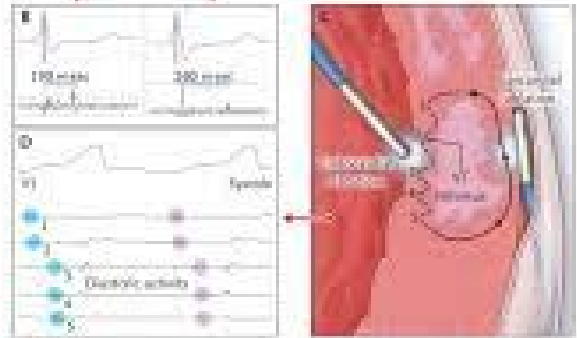
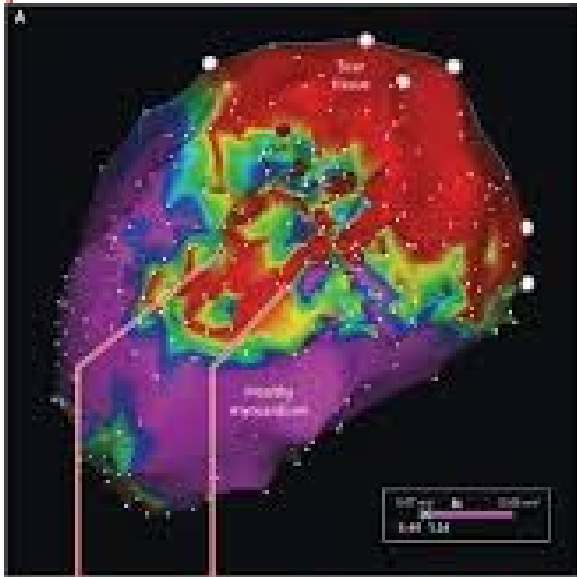
Anti-arrhythmic drug	Effects on ECG	Indications (specific indication)	Oral dose per day (i.v. dose)	Side effects	Contraindications, precautions, other considerations
Amiodarone	Decreases sinus node frequency, prolongs QT interval <sup>a</sup>	PVC, VT, VF	200–400 mg Loading dose: 600–1200 mg/24 h 8–10 days. (Loading dose: 5 mg/kg in 20 min–2 h, 2–3 times in 24 h, then 600–1200 mg/24 h 8–10 days)	<i>Cardiac:</i> Bradycardia, TdP (infrequent) <i>Extracardiac:</i> Photosensitivity, corneal deposits, hypothyroidism, hyperthyroidism, pulmonary toxicity, hepatotoxicity, polyneuropathy, skin discoloration	<i>Precautions:</i> Sinus node dysfunction, severe AV conduction disturbances, hyperthyroidism <i>Other considerations:</i> Can be used in patients with heart failure. Increases the risk of myopathy when used with statins



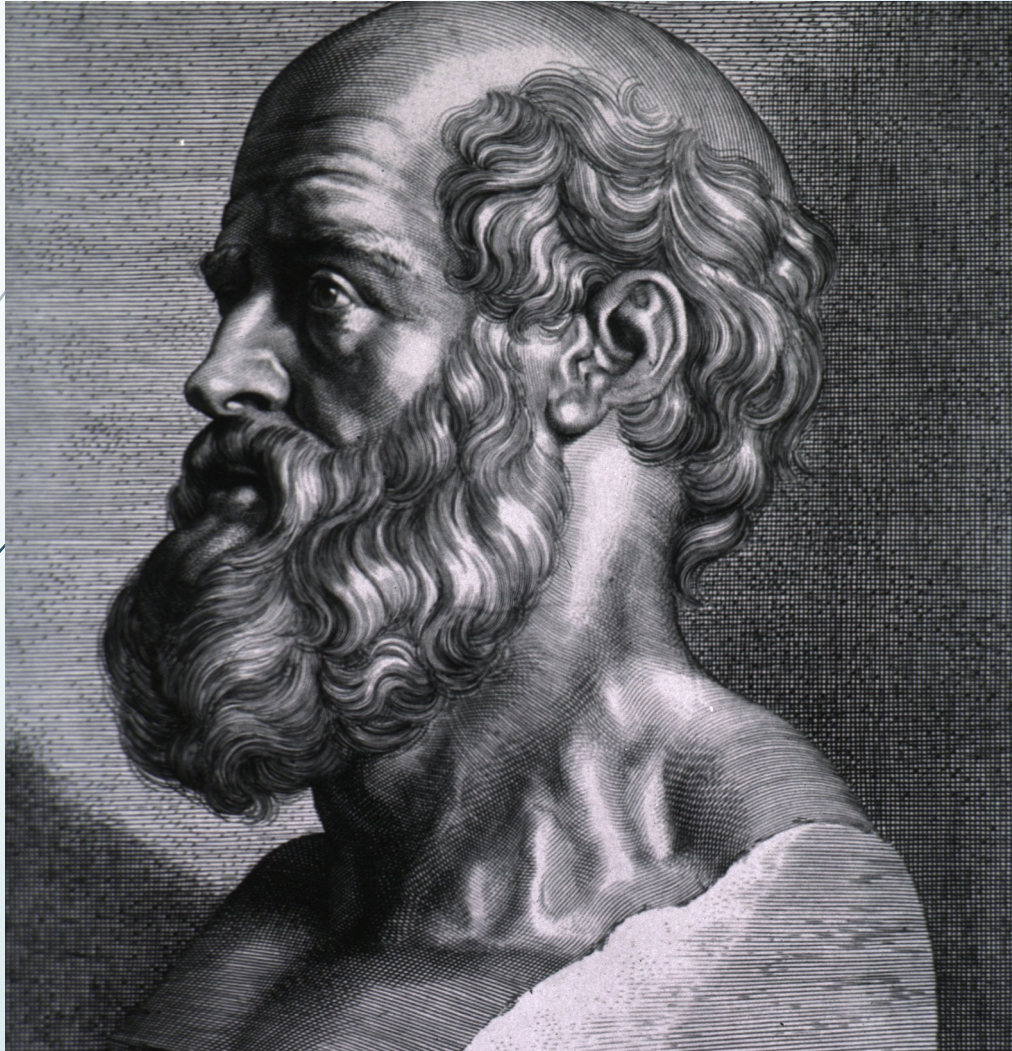
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Anti-arrhythmic drug	Effects on ECG	Indications (specific indication)	Oral dose per day (i.v. dose)	Side effects	Contraindications, precautions, other considerations
Lidocaine	No significant effects	(VT/VF associated with ACS)	No oral use (50–200 mg bolus, then 2–4 mg/min)	<i>Cardiac:</i> Sinoatrial arrest <i>Extracardiac:</i> Central nervous system effects (e.g. drowsiness, dizziness)	<i>Precautions:</i> Reduced dose with reduced liver blood flow (e.g. shock, $\beta$ -blockade, severe heart failure) <i>Other considerations:</i> More effective with high potassium level. Few haemodynamic side effects









Prevent

Cure

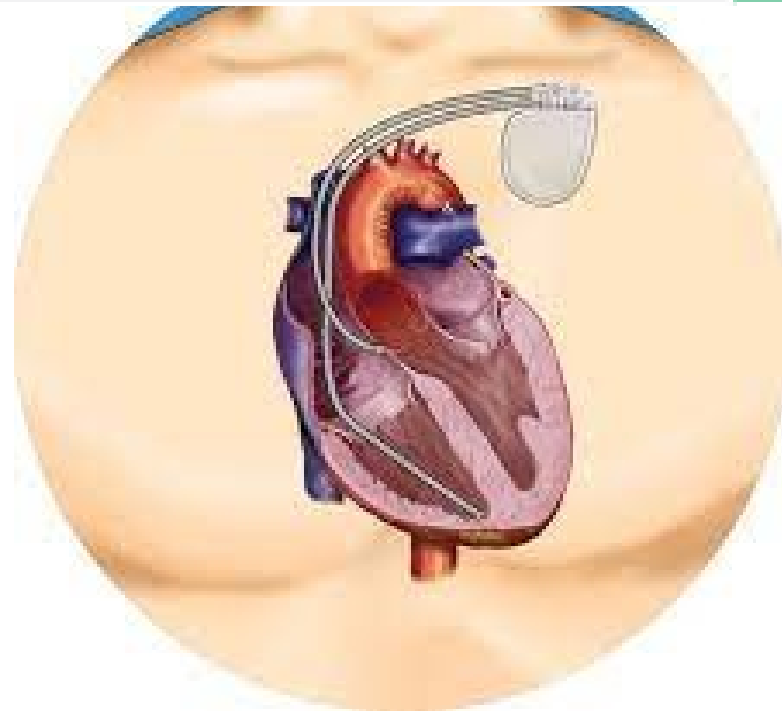


## Secondary prevention of SCD and treatment of VAs

ICD implantation is recommended in patients without ongoing ischaemia with documented VF or haemodynamically not-tolerated VT occurring later than 48 h after MI.<sup>349–351</sup>

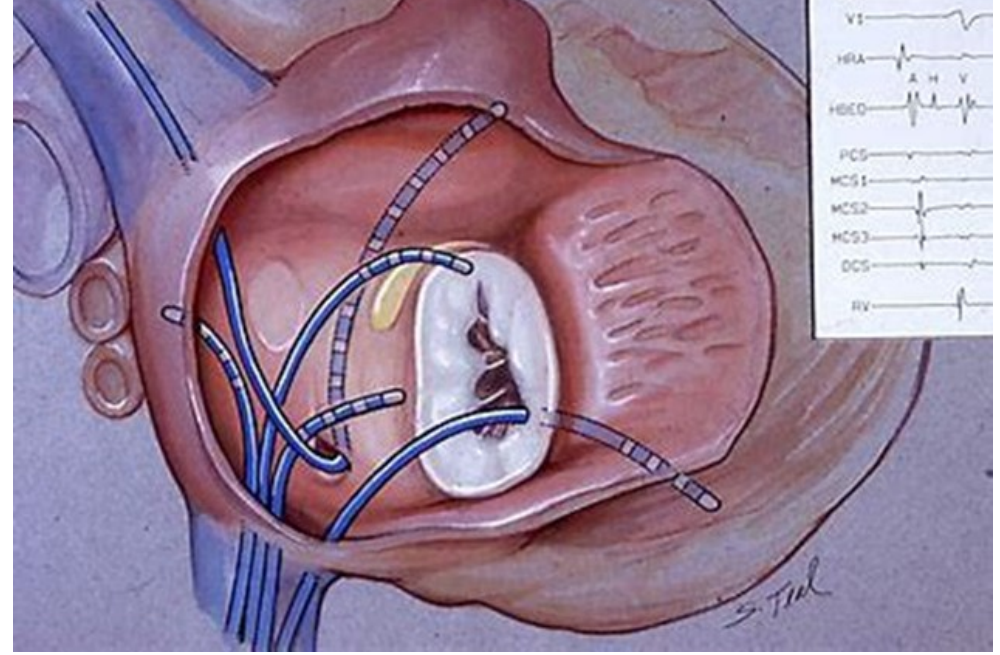
I

A



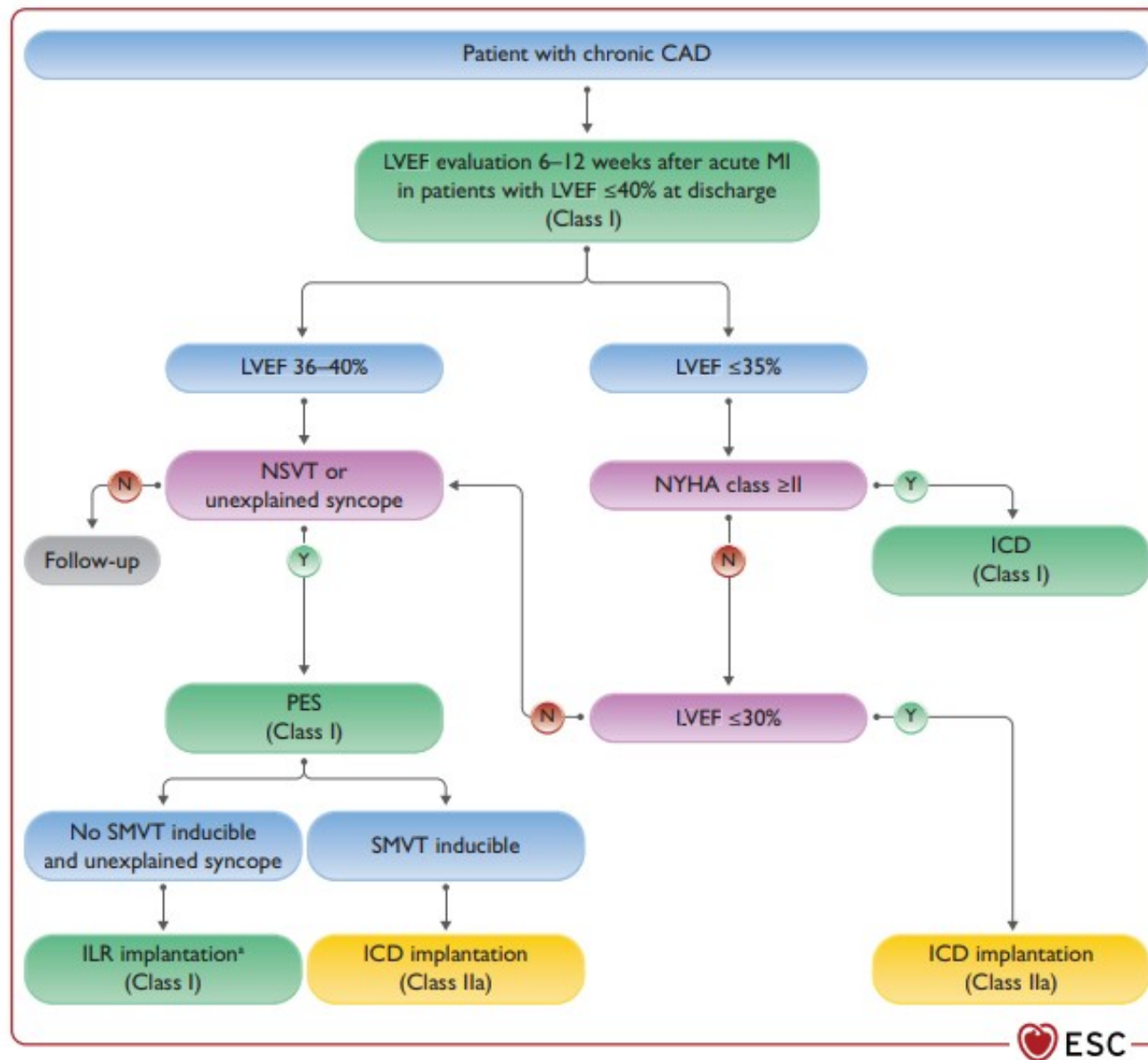
**Recommendation Table 23** — Recommendations for risk stratification and treatment of ventricular arrhythmias early after myocardial infarction

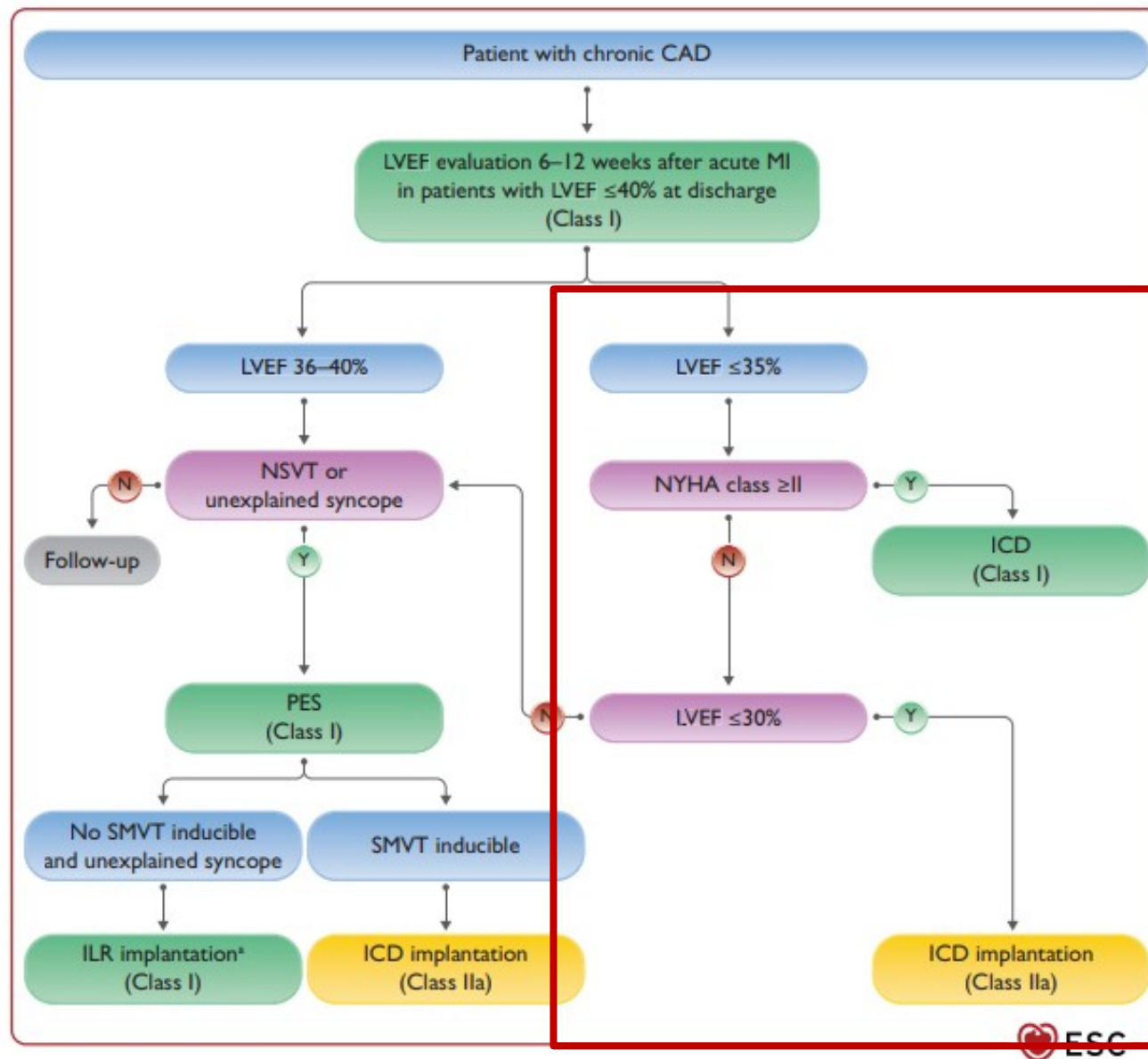
Recommendations	Class <sup>a</sup>	Level <sup>b</sup>
<b>Risk stratification</b>		
Early (before discharge) assessment of LVEF is recommended in all patients with acute MI. <sup>567,568</sup>	<b>I</b>	<b>B</b>
In patients with pre-discharge LVEF ≤40%, re-evaluation of LVEF 6–12 weeks after MI is recommended to assess the potential need for primary prevention ICD implantation. <sup>568,573,574</sup>	<b>I</b>	<b>C</b>

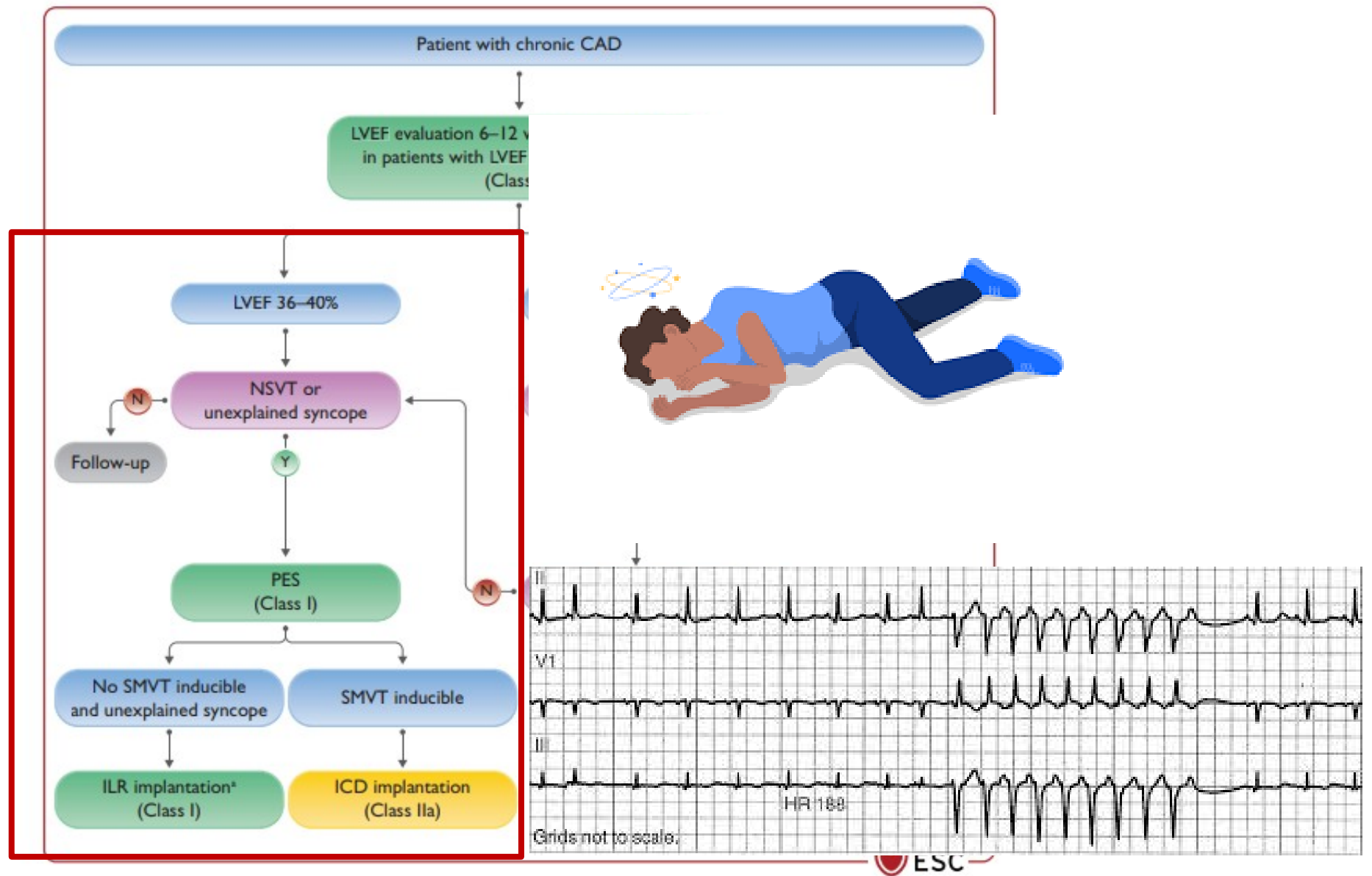


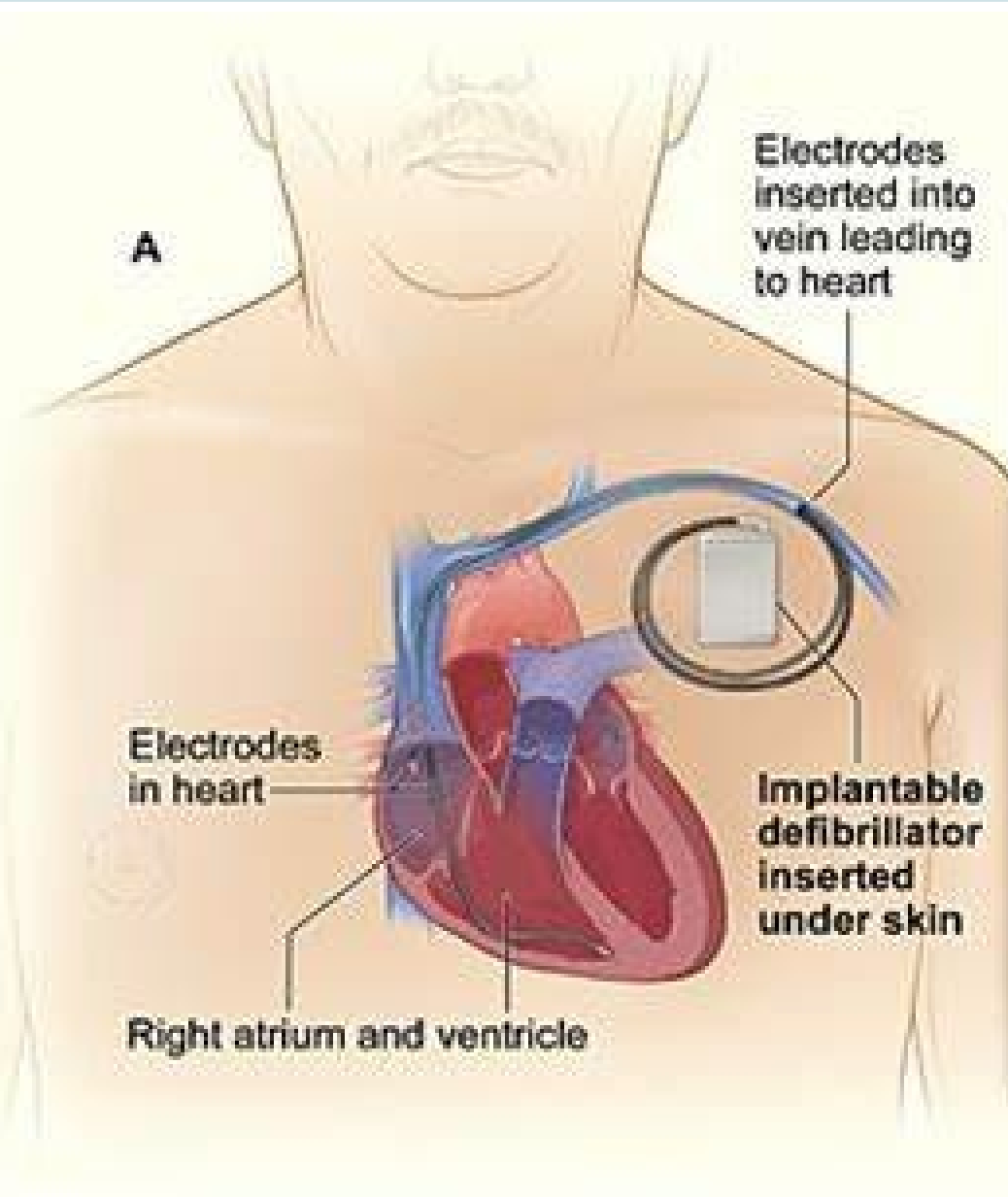
Εργαλεία











**A**

**Electrodes  
inserted into  
vein leading  
to heart**

**Electrodes  
in heart**

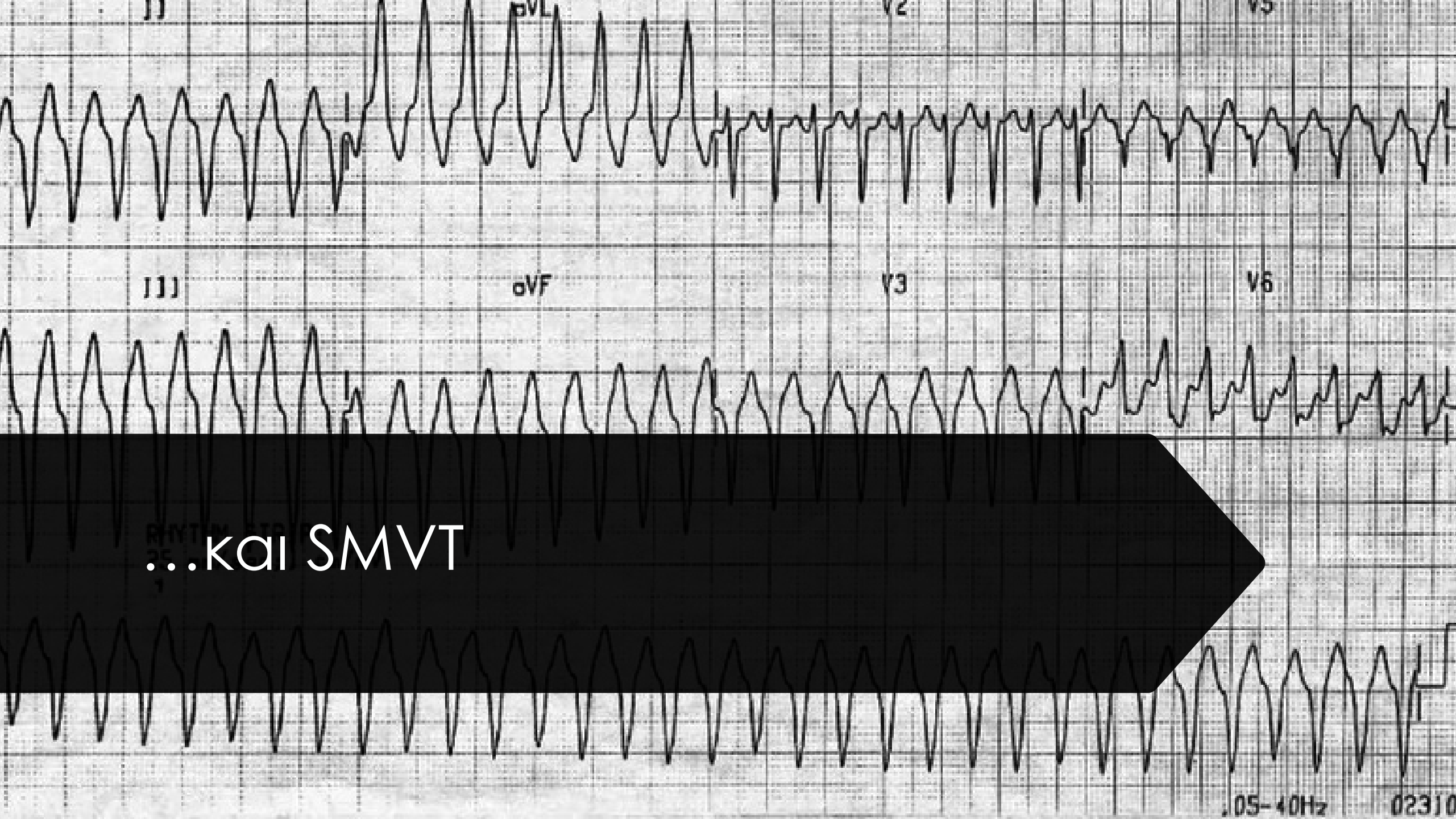
**Implantable  
defibrillator  
inserted  
under skin**

**Right atrium and ventricle**

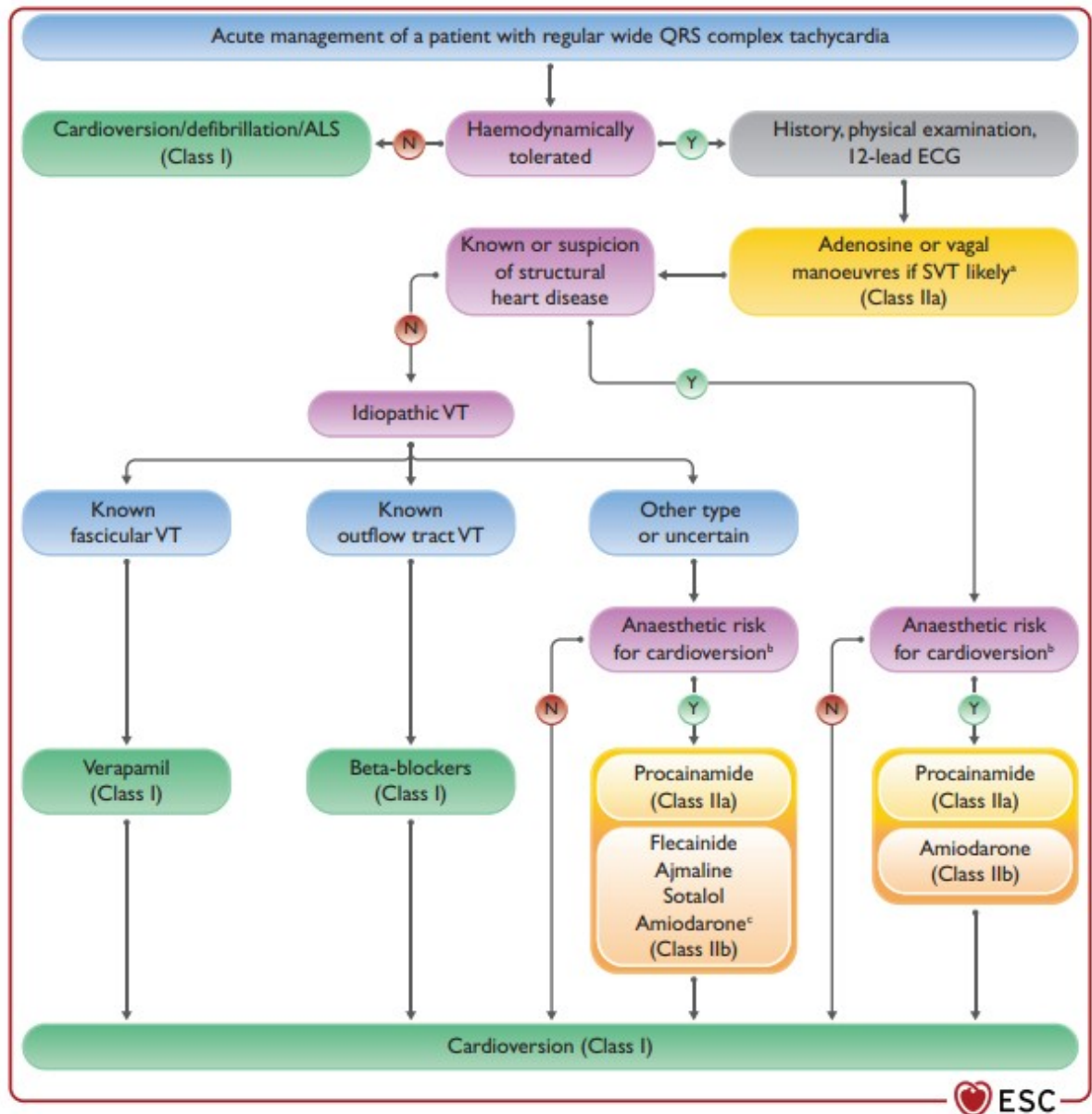
Ασθενής με ιστορικό στεφανιαίας νόσου...

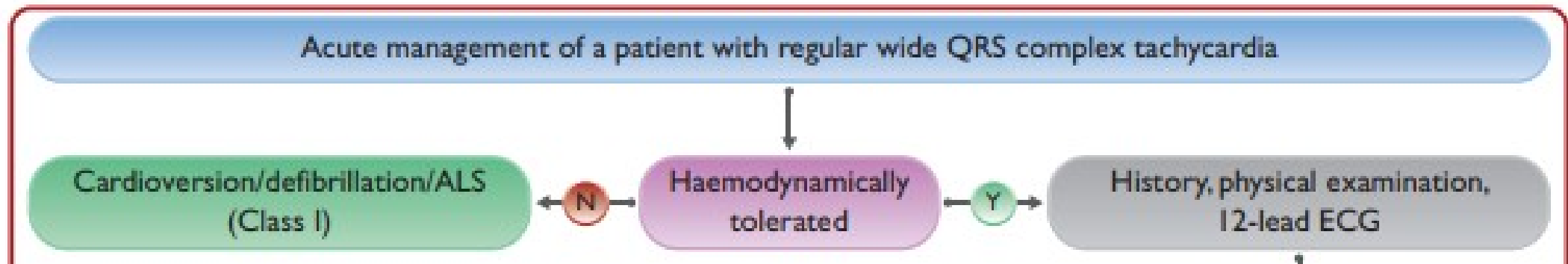
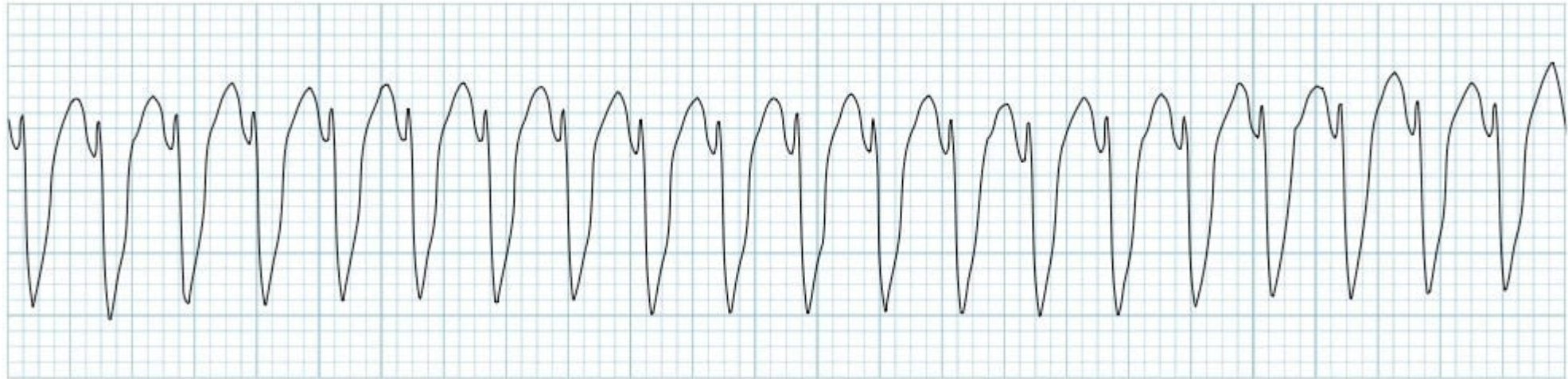


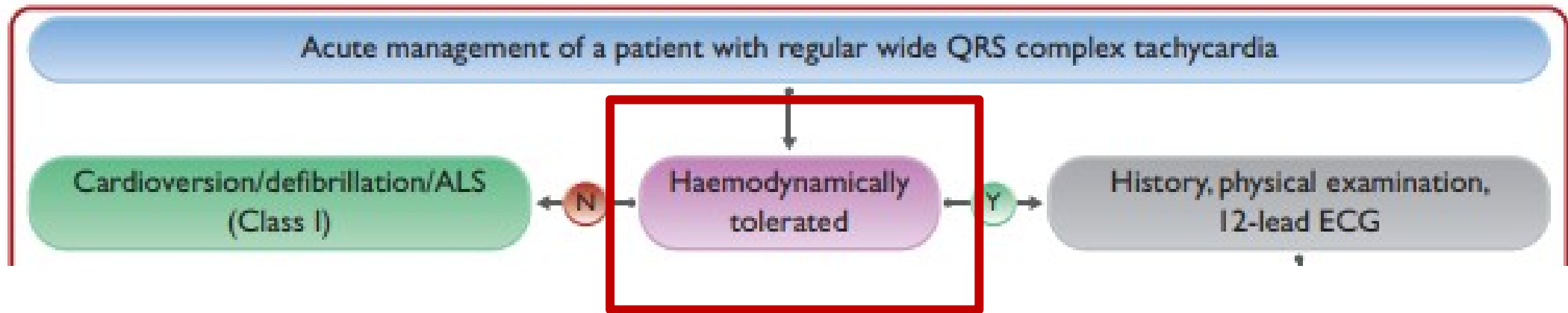
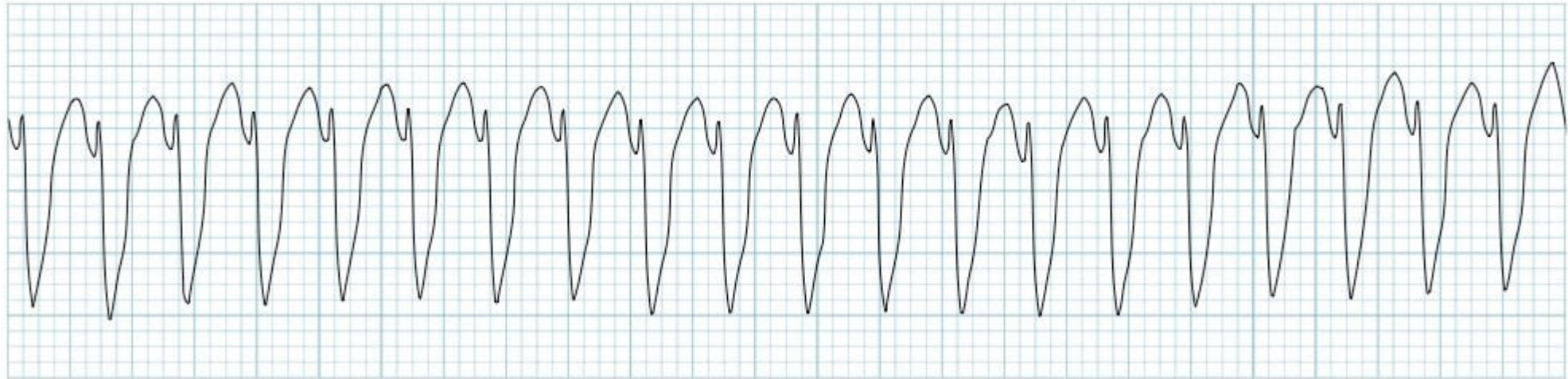


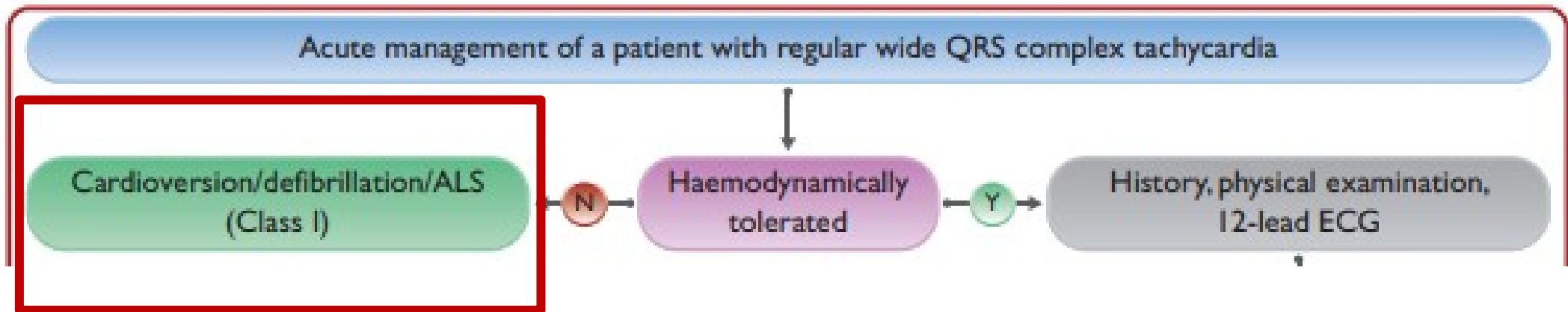
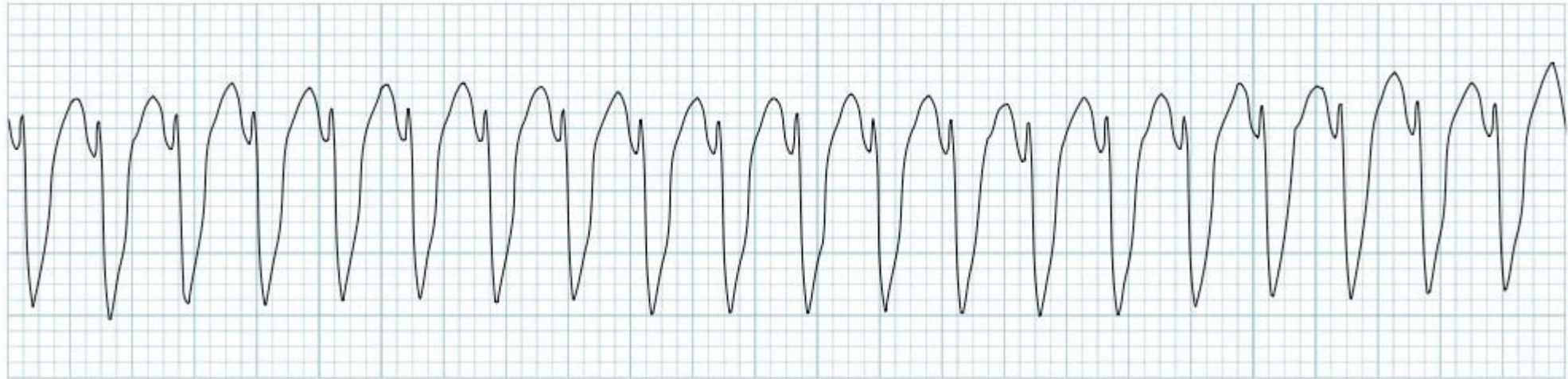


RHYTHM: SINOBRADYCARDIA  
...kai SMVT

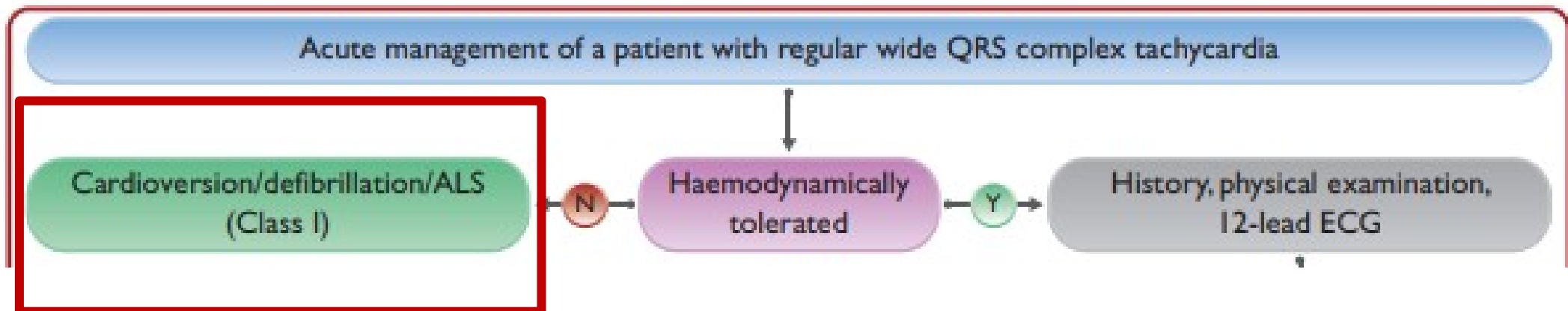


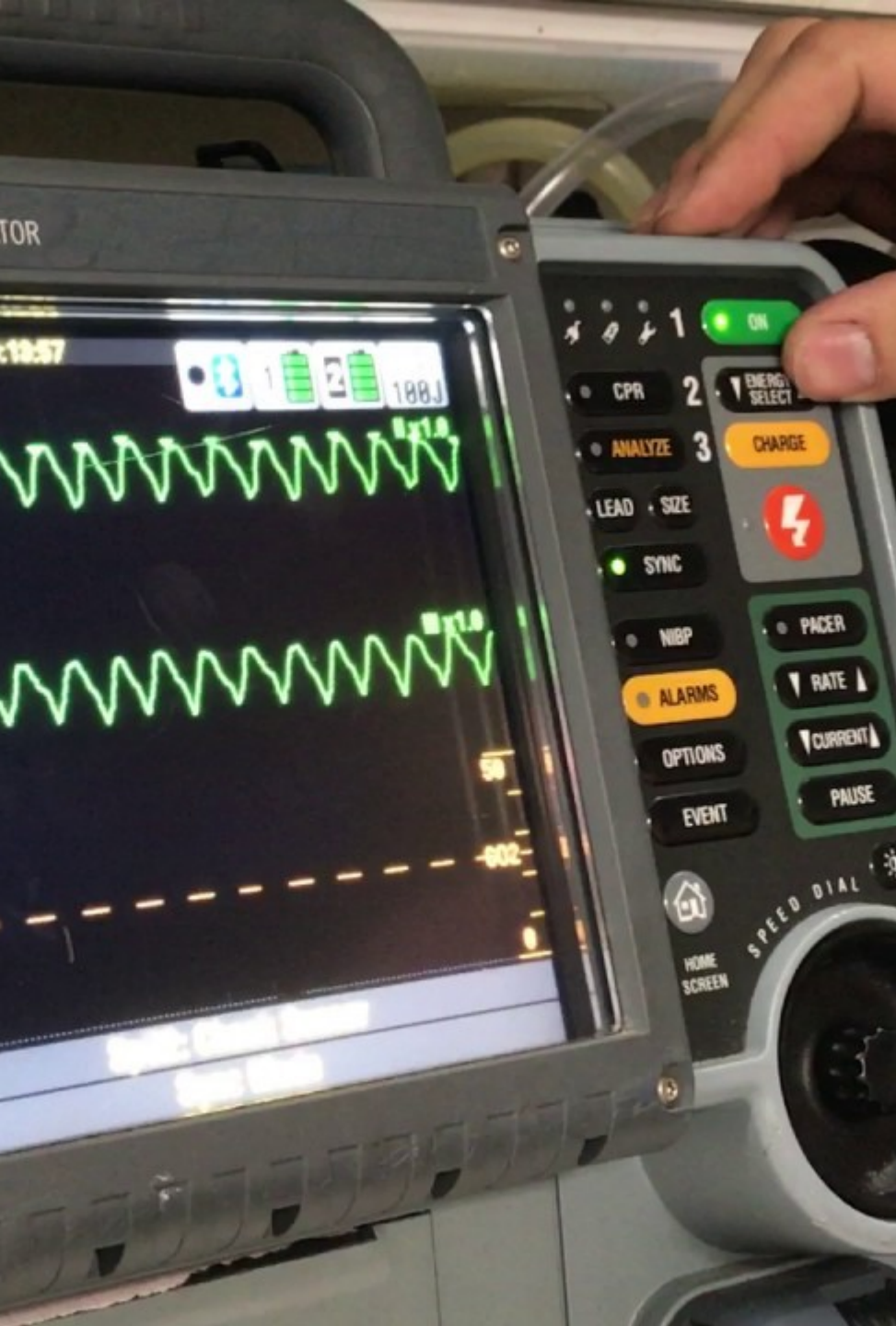






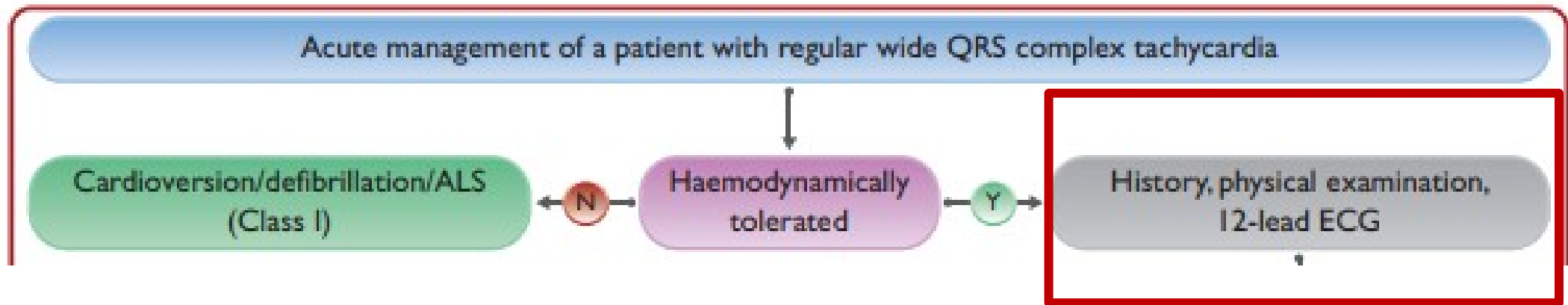
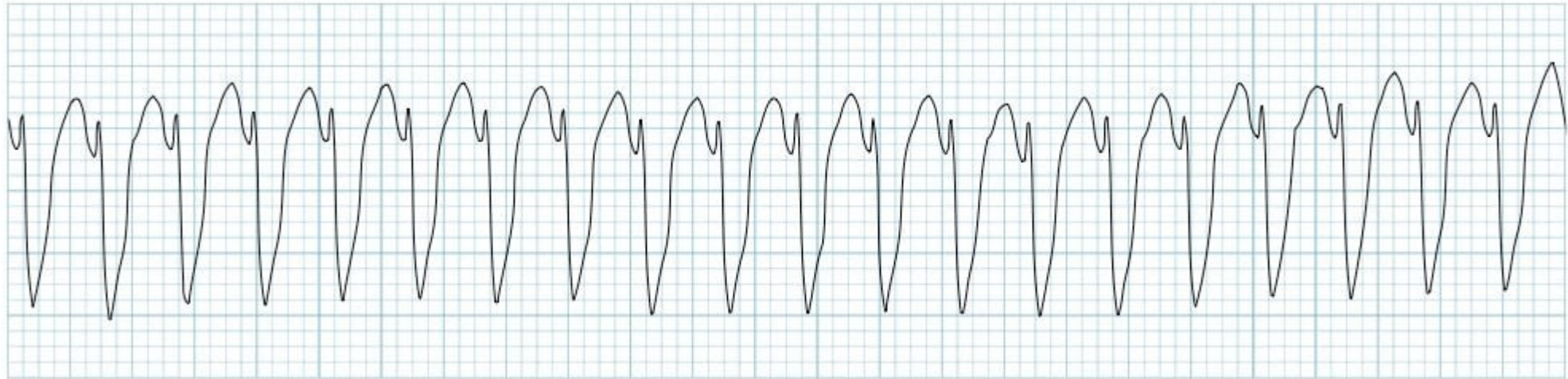
Recommendations	Class <sup>a</sup>	Level <sup>b</sup>
<b>Acute management of sustained VT</b>		
DC cardioversion is recommended as the first-line treatment for patients with haemodynamically not-tolerated SMVT. <sup>303,339</sup>	<b>I</b>	<b>B</b>



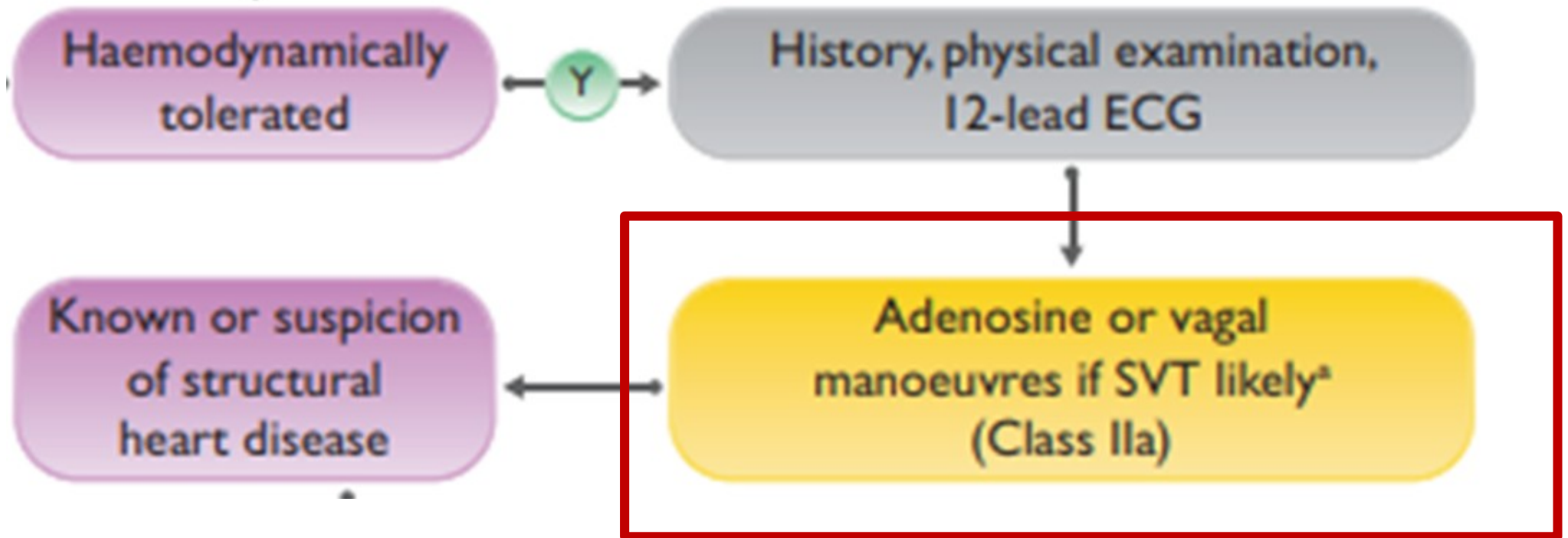


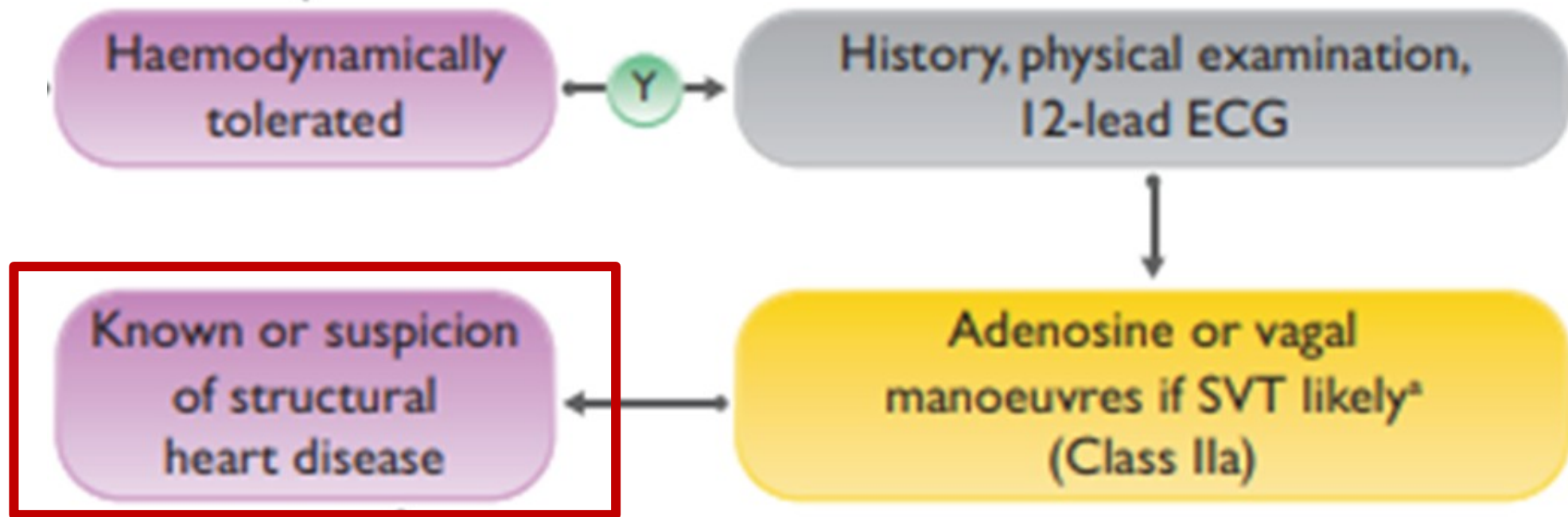
## Ηλεκτρική καρδιομετατροπή- απινίδωση

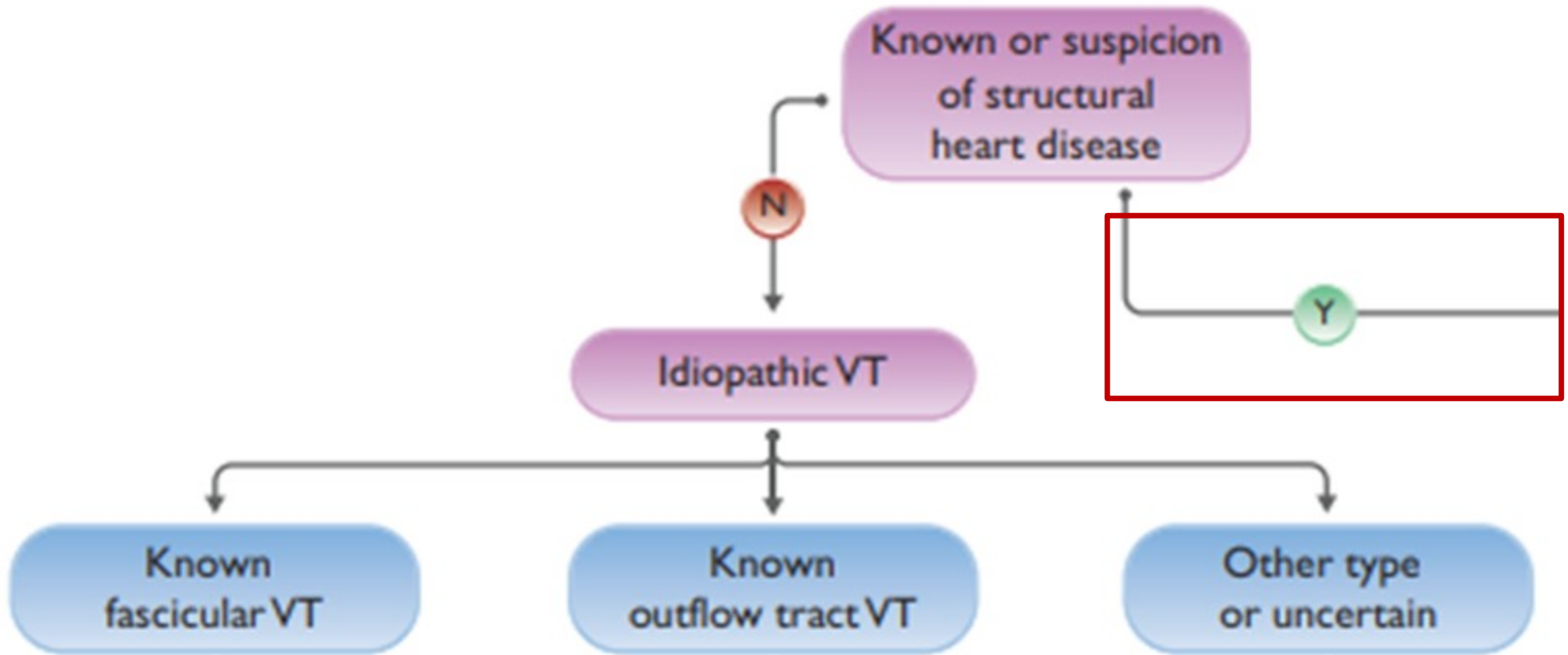
- Συγχρονισμένη καρδιομετατροπή
- Εάν ο συγχρονισμός δεν είναι εφικτός, ασύγχρονη χορήγηση σοκ





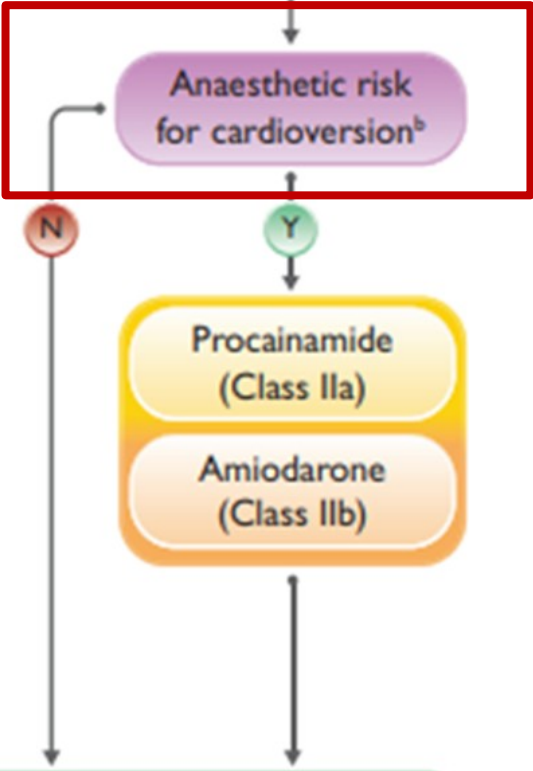




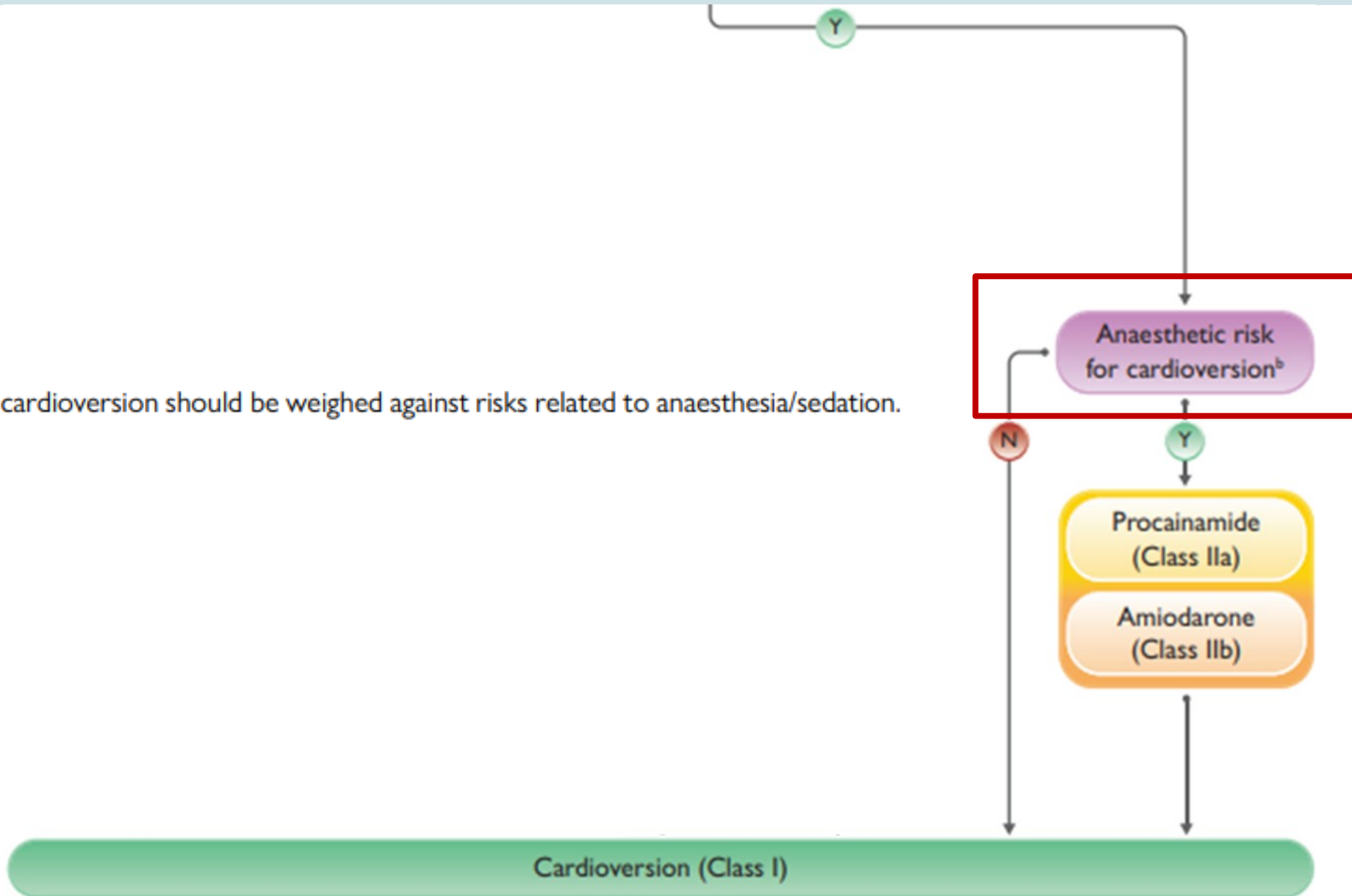


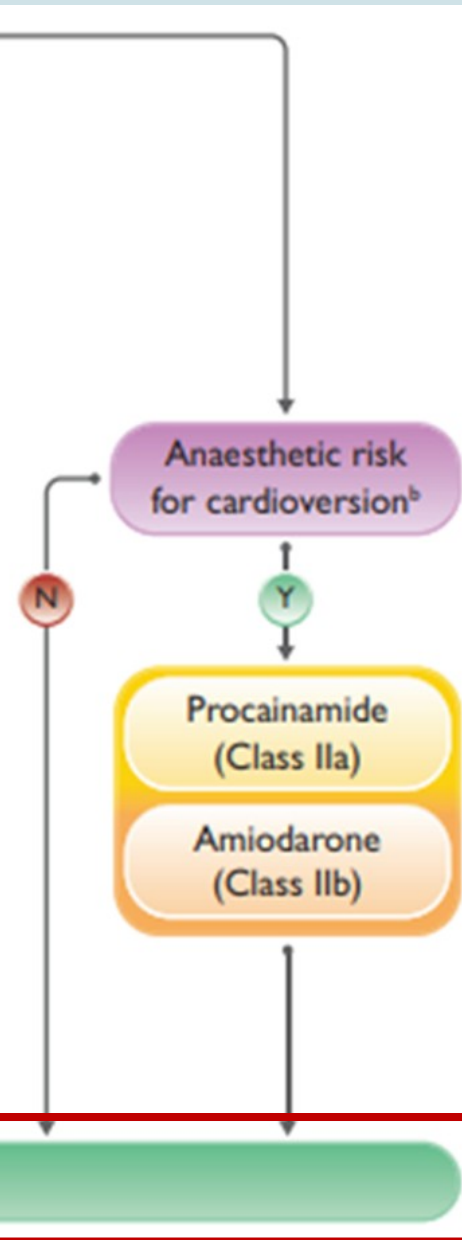


Cardioversion (Class I)



<sup>b</sup>Benefit of cardioversion should be weighed against risks related to anaesthesia/sedation.



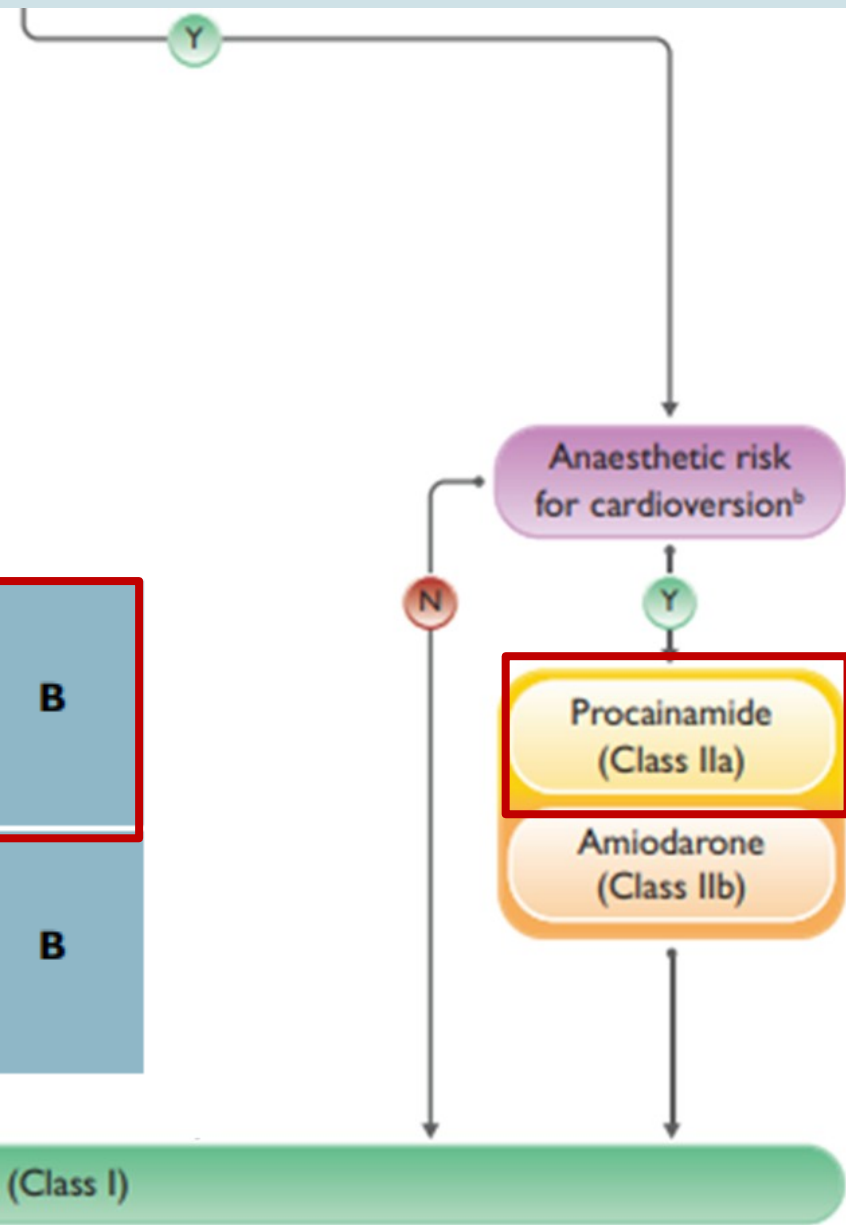


Cardioversion (Class I)



In patients presenting with a haemodynamically tolerated SMVT and known or suspected SHD, intravenous procainamide should be considered. <sup>303</sup>	<b>IIa</b>	<b>B</b>
In patients presenting with a haemodynamically tolerated SMVT in the absence of an established diagnosis, intravenous amiodarone may be considered. <sup>303</sup>	<b>IIb</b>	<b>B</b>

Cardioversion (Class I)



**Table 8** Anti-arrhythmic drugs (acute and chronic treatment)

Anti-arrhythmic drug	Effects on ECG	Indications (specific indication)	Oral dose per day (i.v. dose)	Side effects	Contraindications, precautions, other considerations
Procainamide	Prolongs PR interval, QRS duration, and QT interval <sup>a</sup>	VT	(100 mg bolus, can be repeated after 5 min if no effect, max 500–750 mg [max 50 mg/min]. Then, 2–6 mg/min)	<i>Cardiac:</i> Sinus bradycardia, hypotension, TdP <i>Extracardiac:</i> Rash, myalgia, vasculitis, systemic lupus, agranulocytosis	<i>Contraindications:</i> Severe sinus node dysfunction, severe AV conduction disturbances, severe intraventricular conduction disturbances, severe LV dysfunction hypotension, BrS





In patients presenting with a haemodynamically tolerated SMVT and known or suspected SHD, intravenous procainamide should be considered.<sup>303</sup>

In patients presenting with a haemodynamically tolerated SMVT in the absence of an established diagnosis, intravenous amiodarone may be considered.<sup>303</sup>

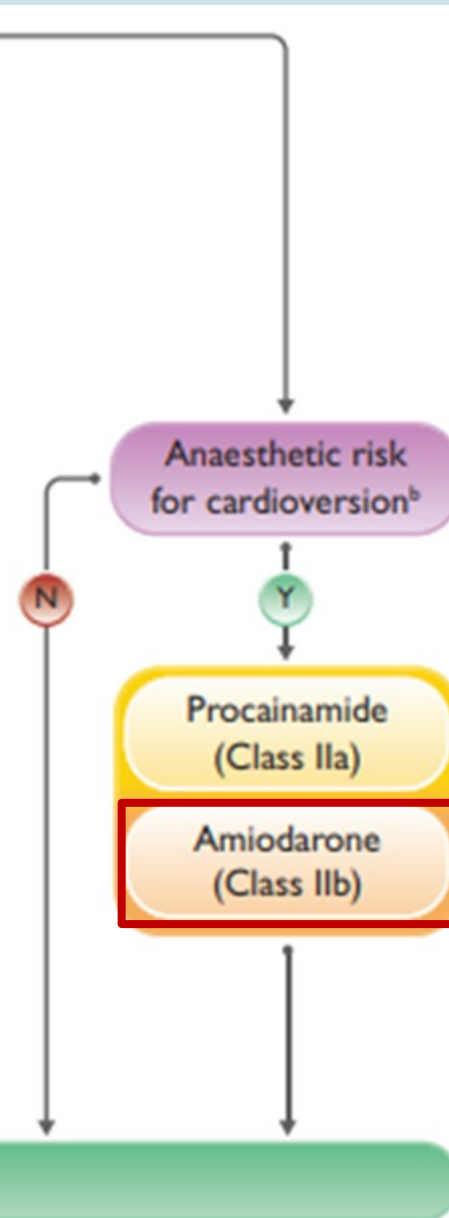
**IIa**

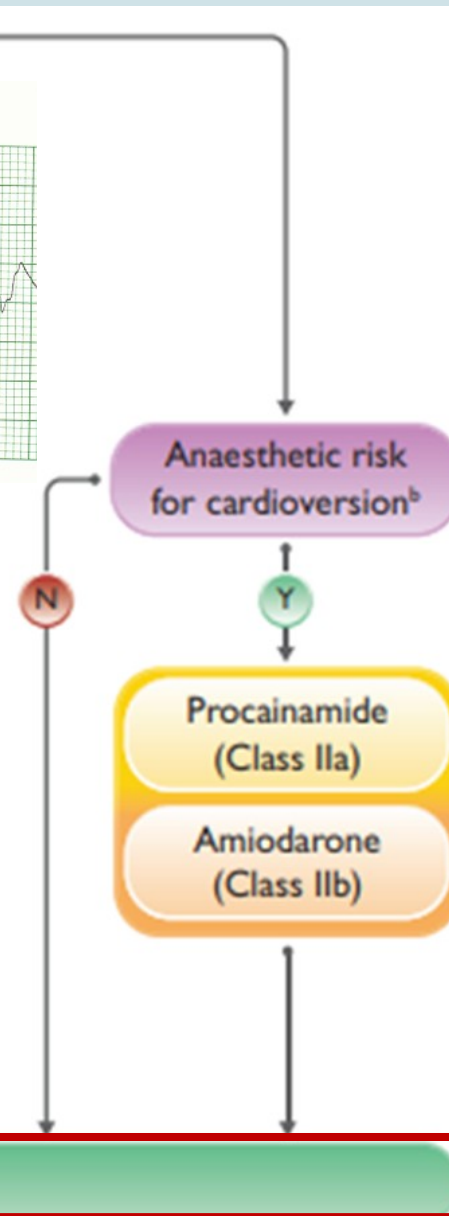
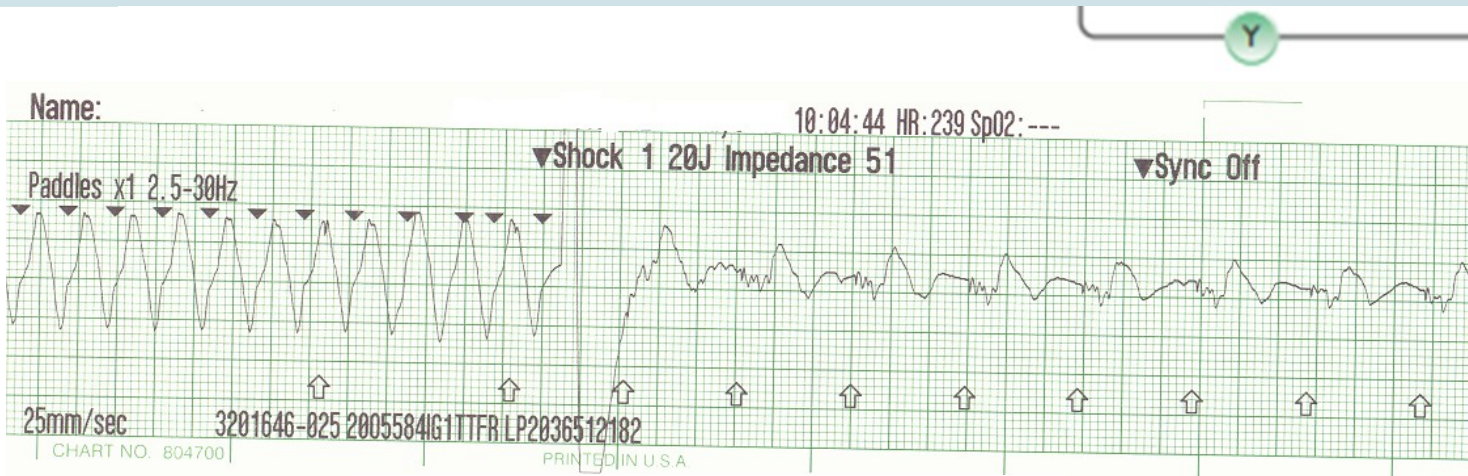
**B**

**IIb**

**B**

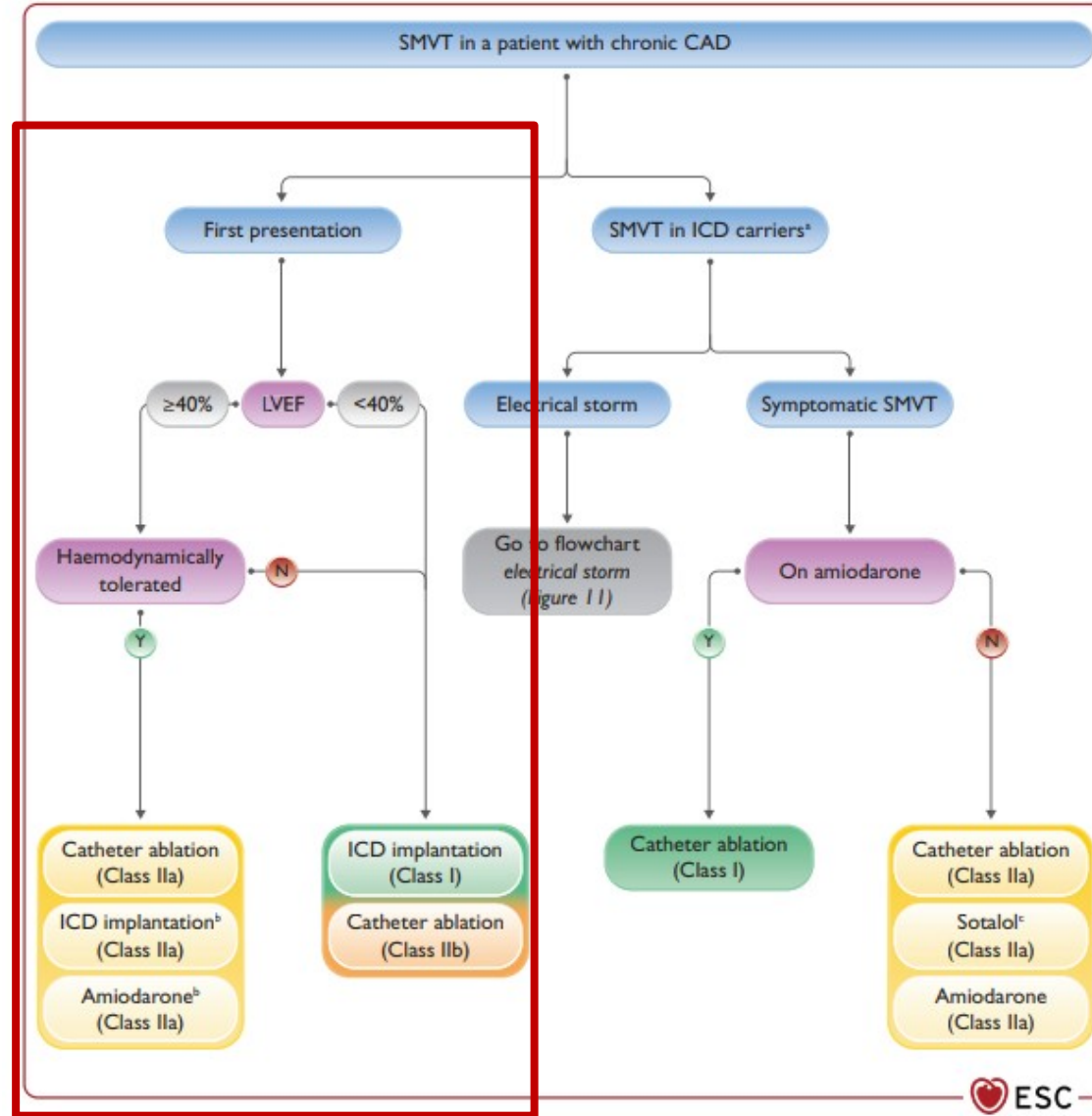
Cardioversion (Class I)

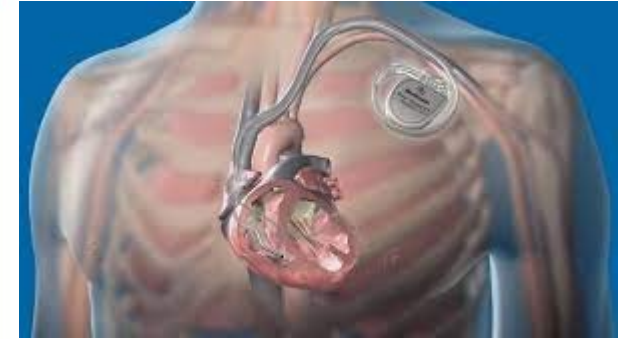
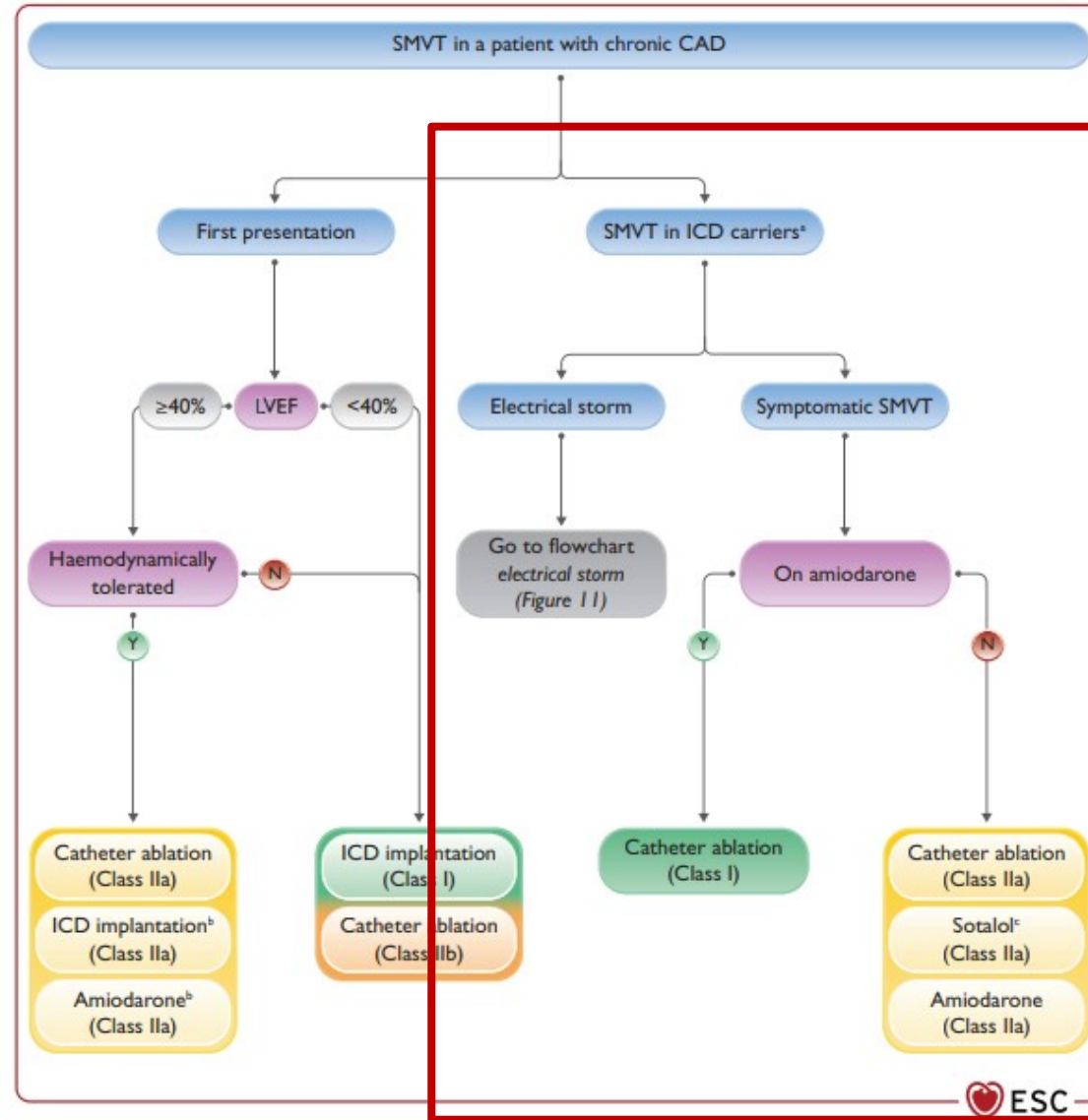




Cardioversion (Class I)

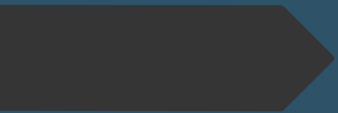


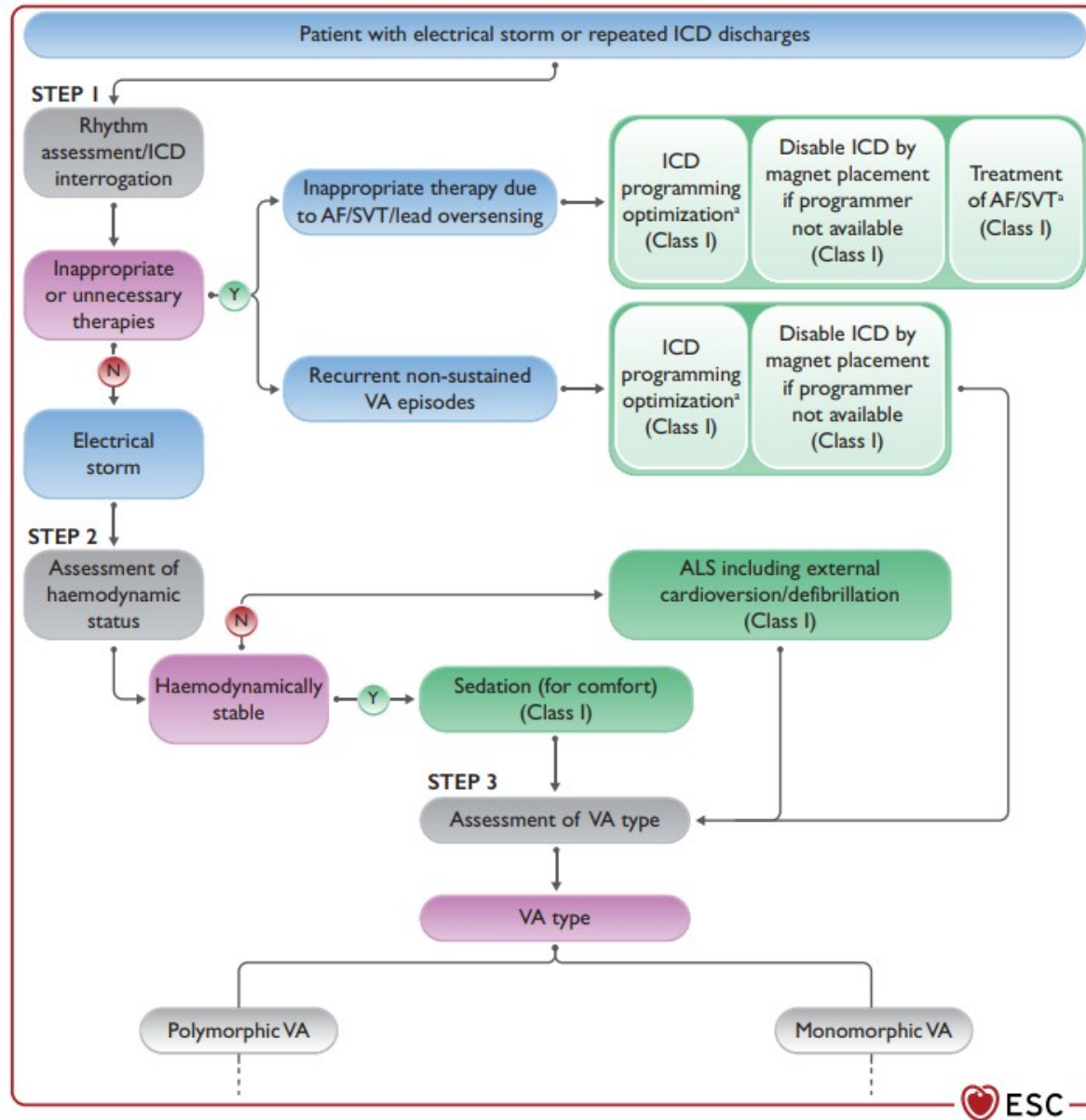


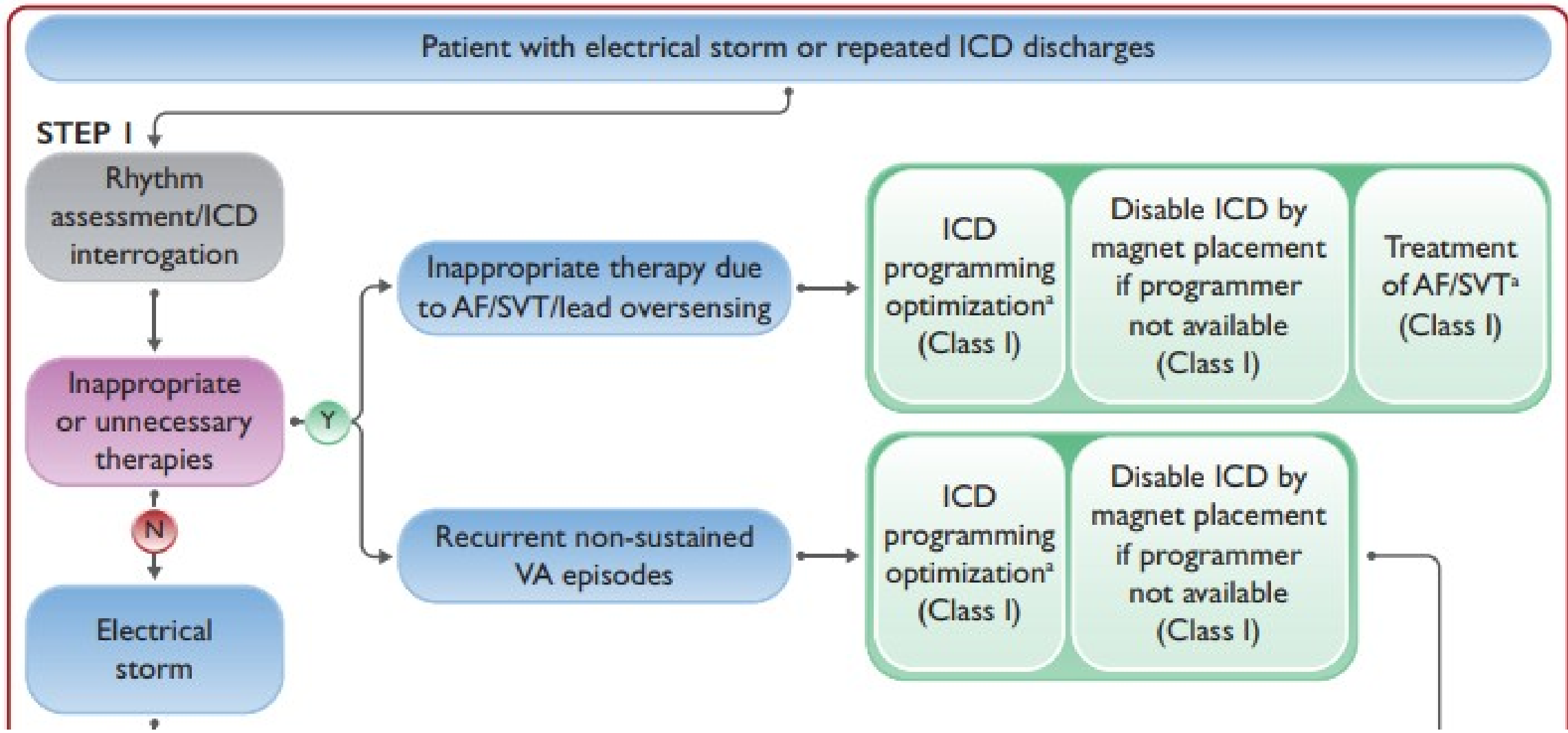


Ασθενής με εμφυτευμένο ICD και...

... ηλεκτρική θύελλα









Patient with electrical storm or repeated ICD discharges

STEP I

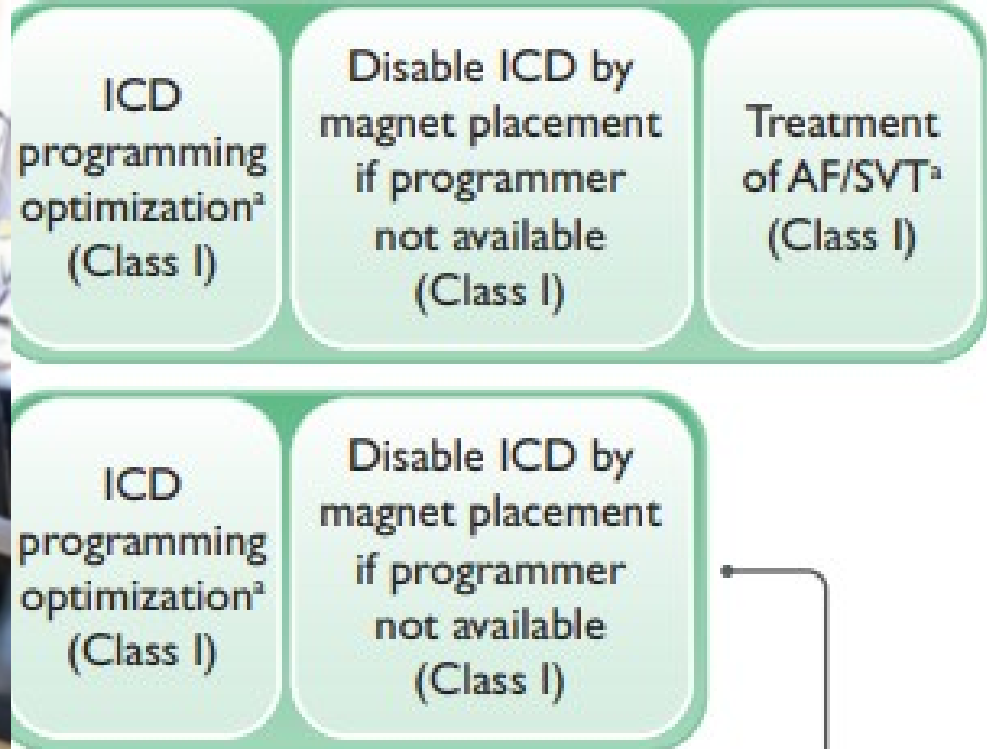
Rhythm assessment/ICD interrogation

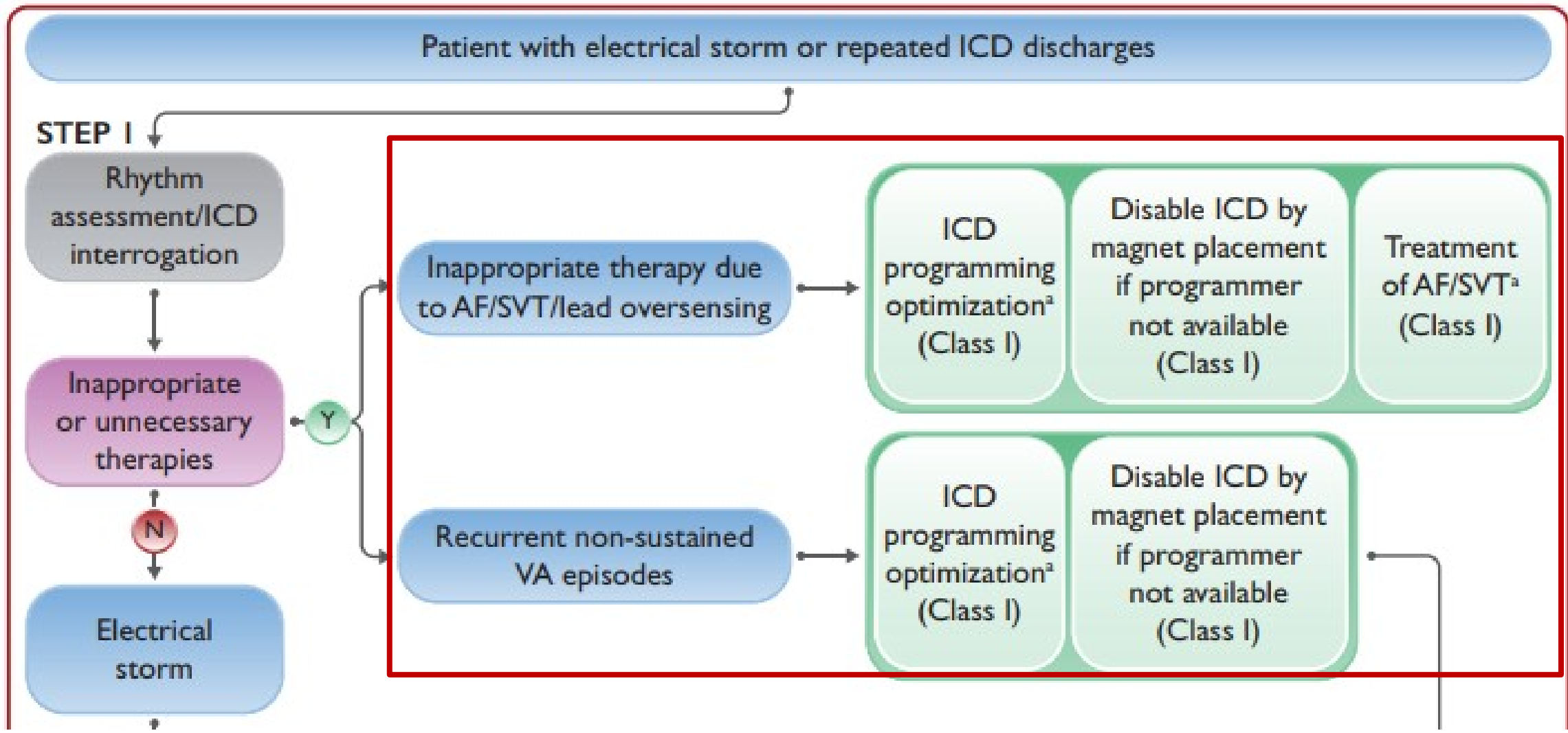
Inappropriate or unnecessary therapies

N

Electrical storm

Y





## Patient with electrical storm or repeated ICD discharges

STEP I

Rhythm  
assessment/ICD  
interrogation

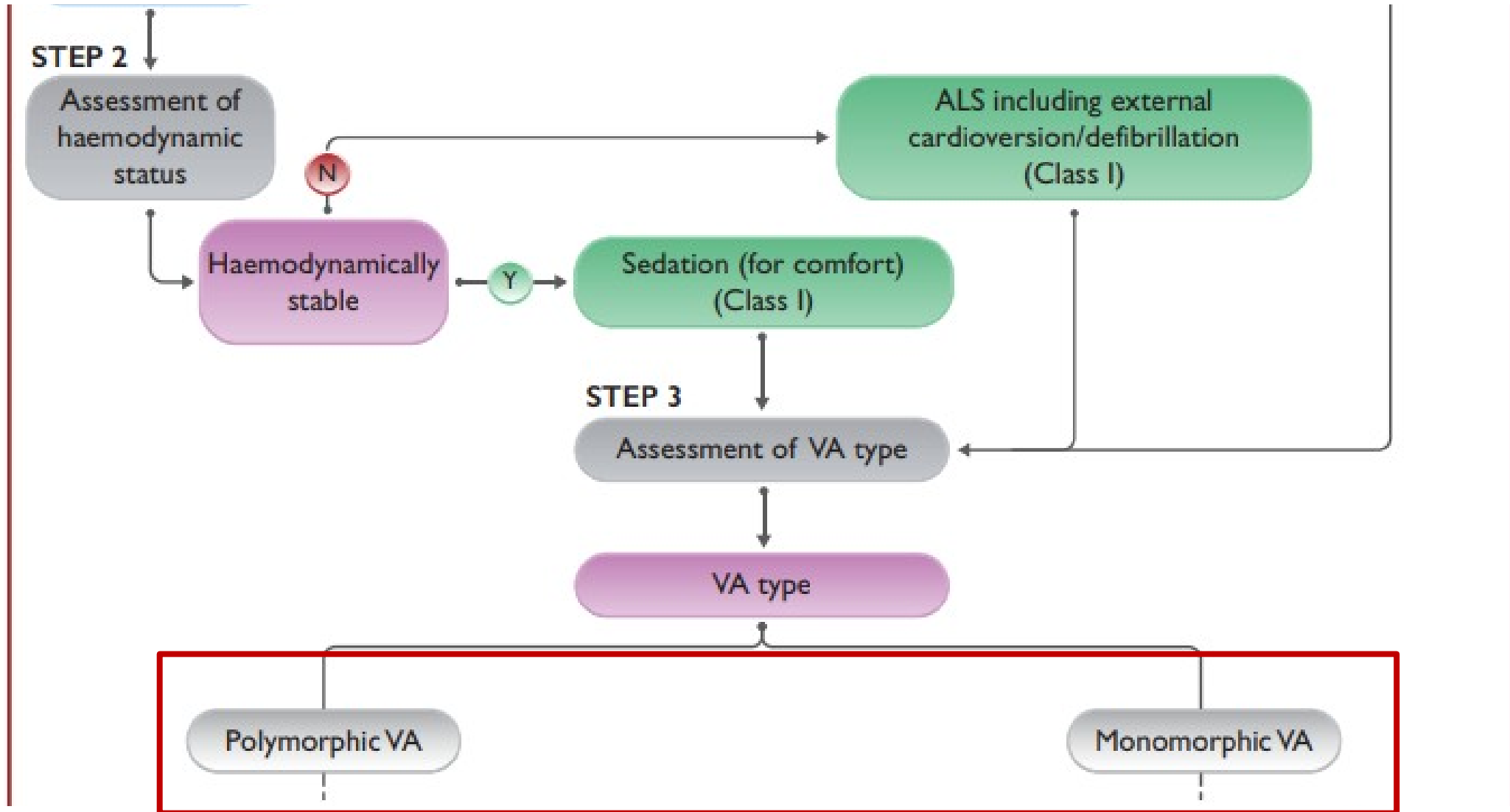
Inappropriate  
or unnecessary  
therapies

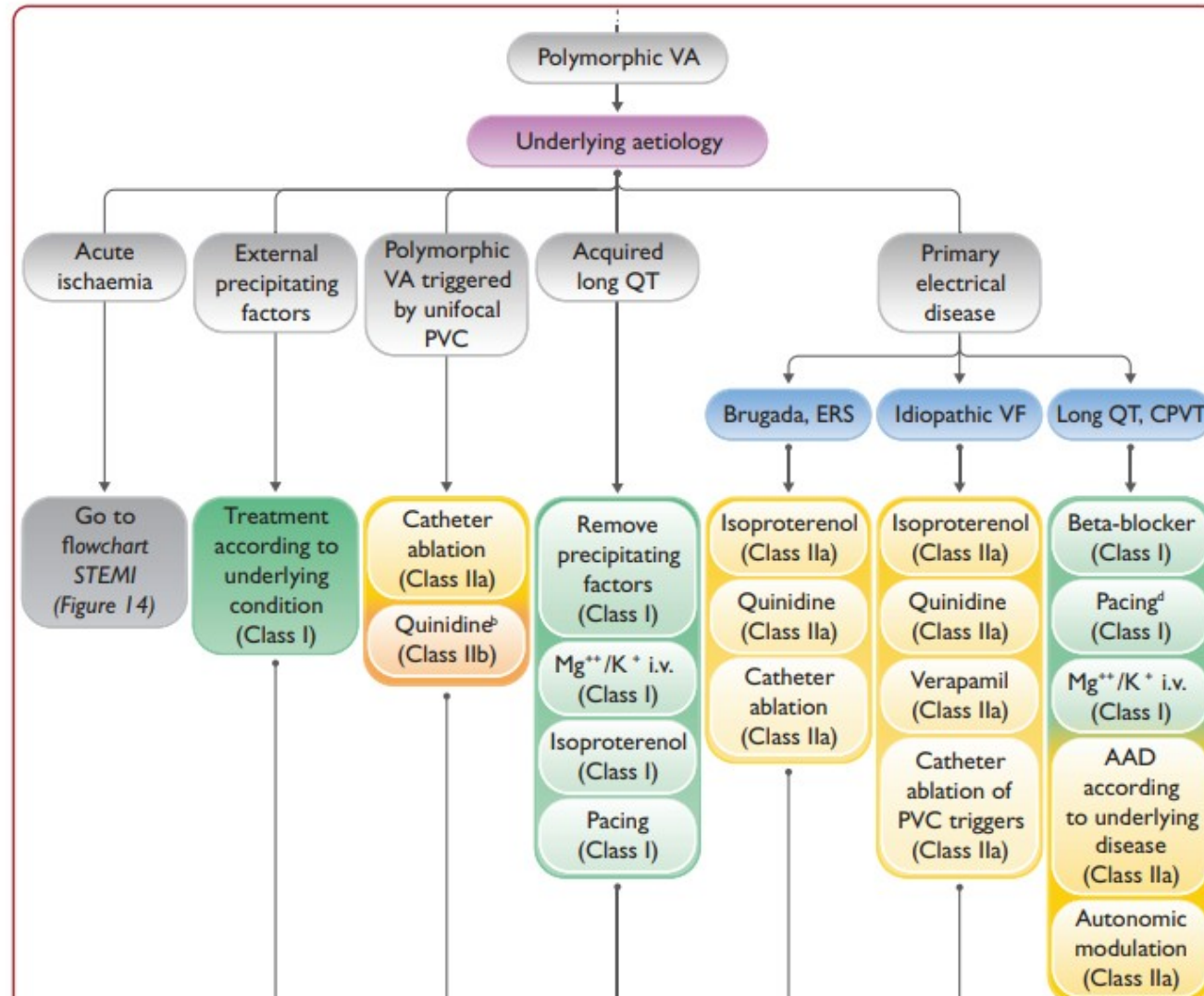
Y

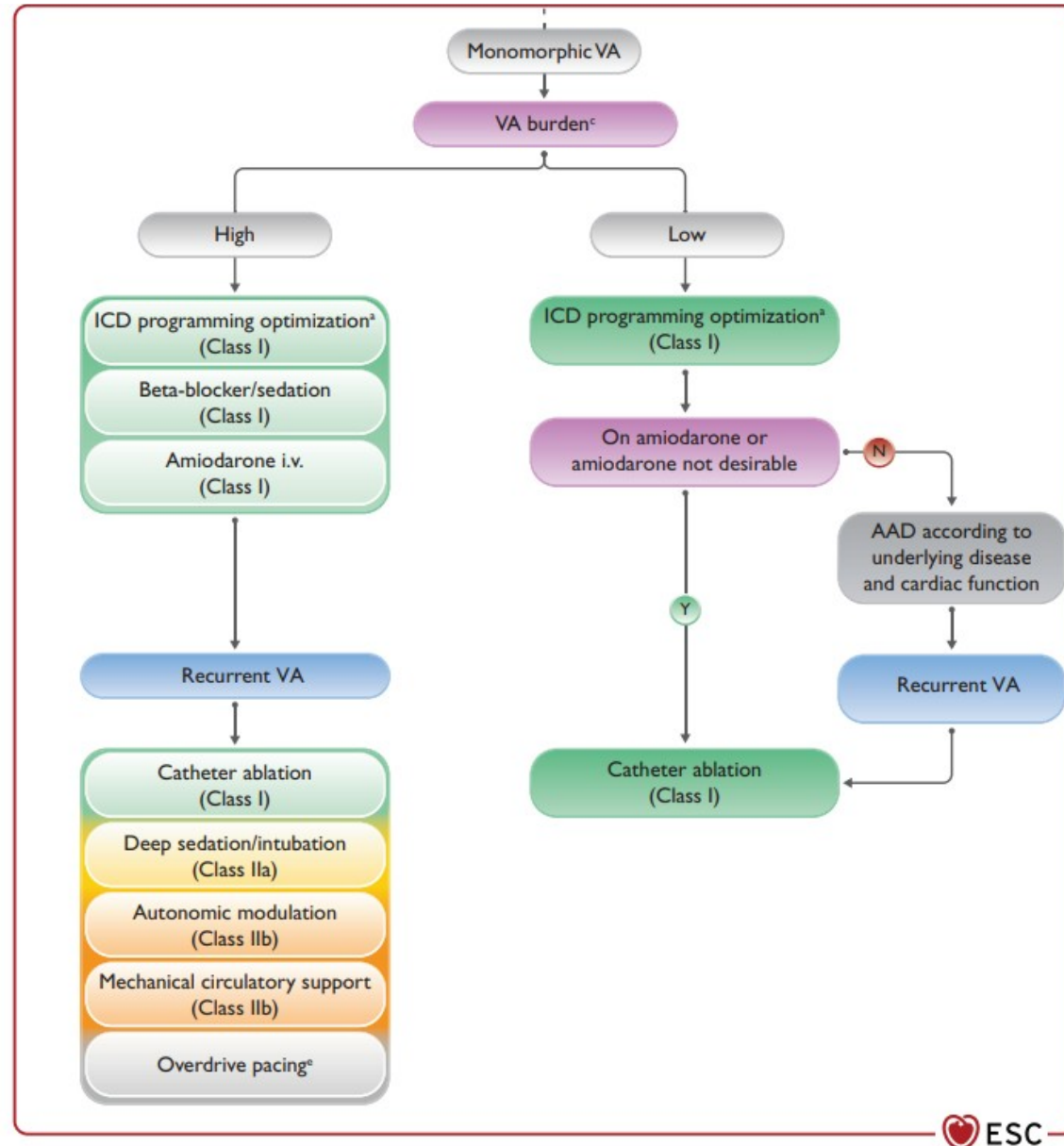
N

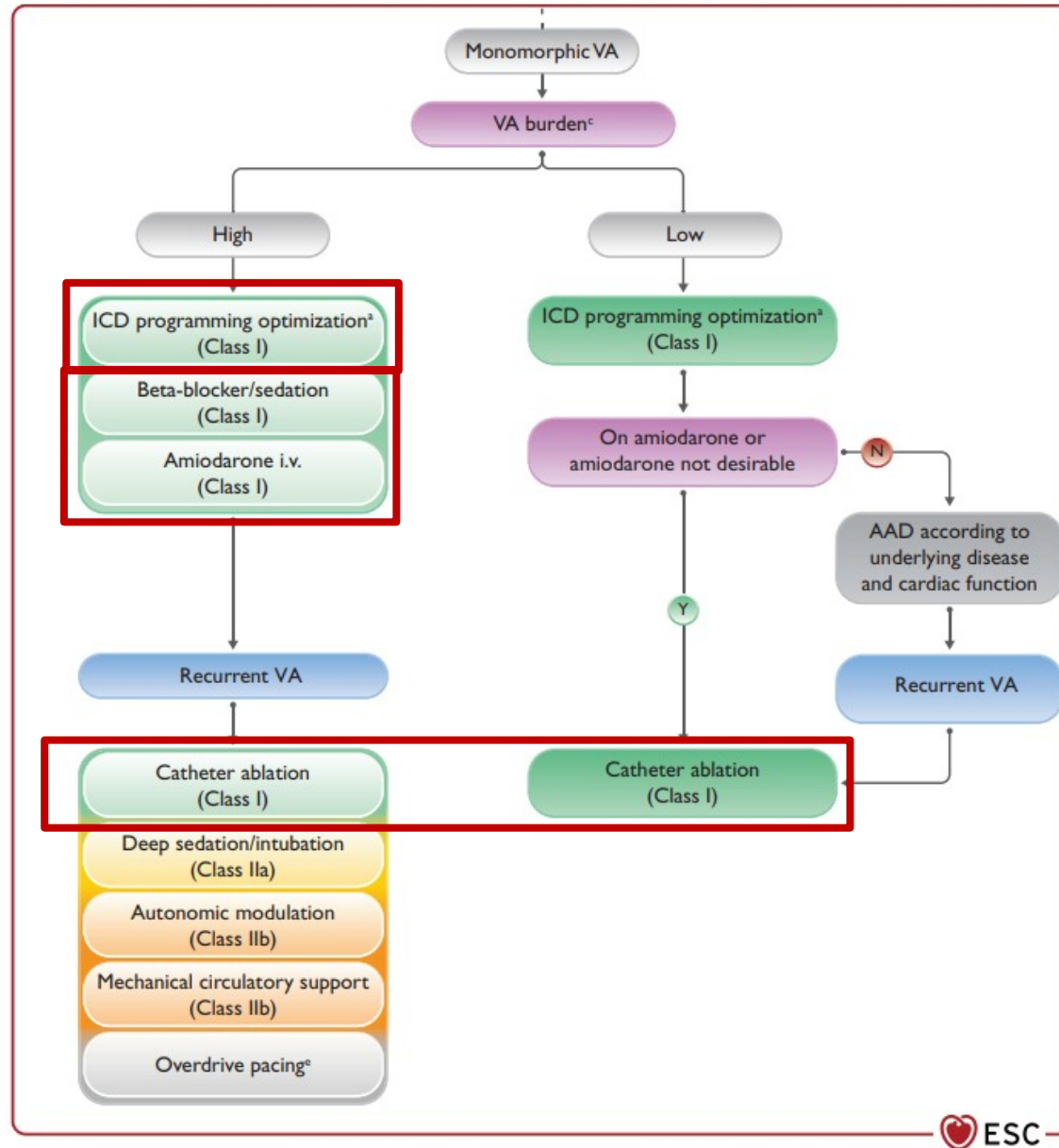
Electrical  
storm







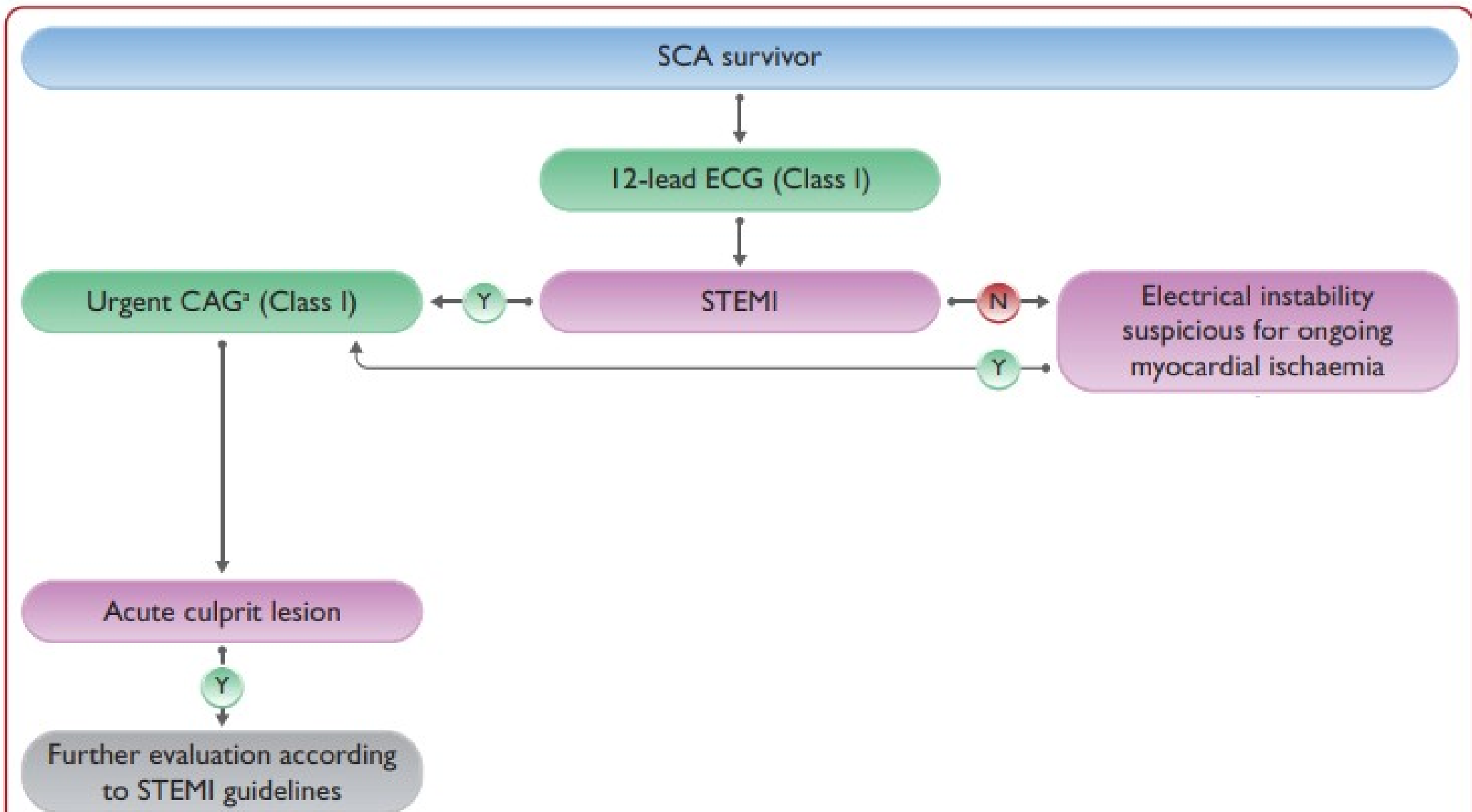


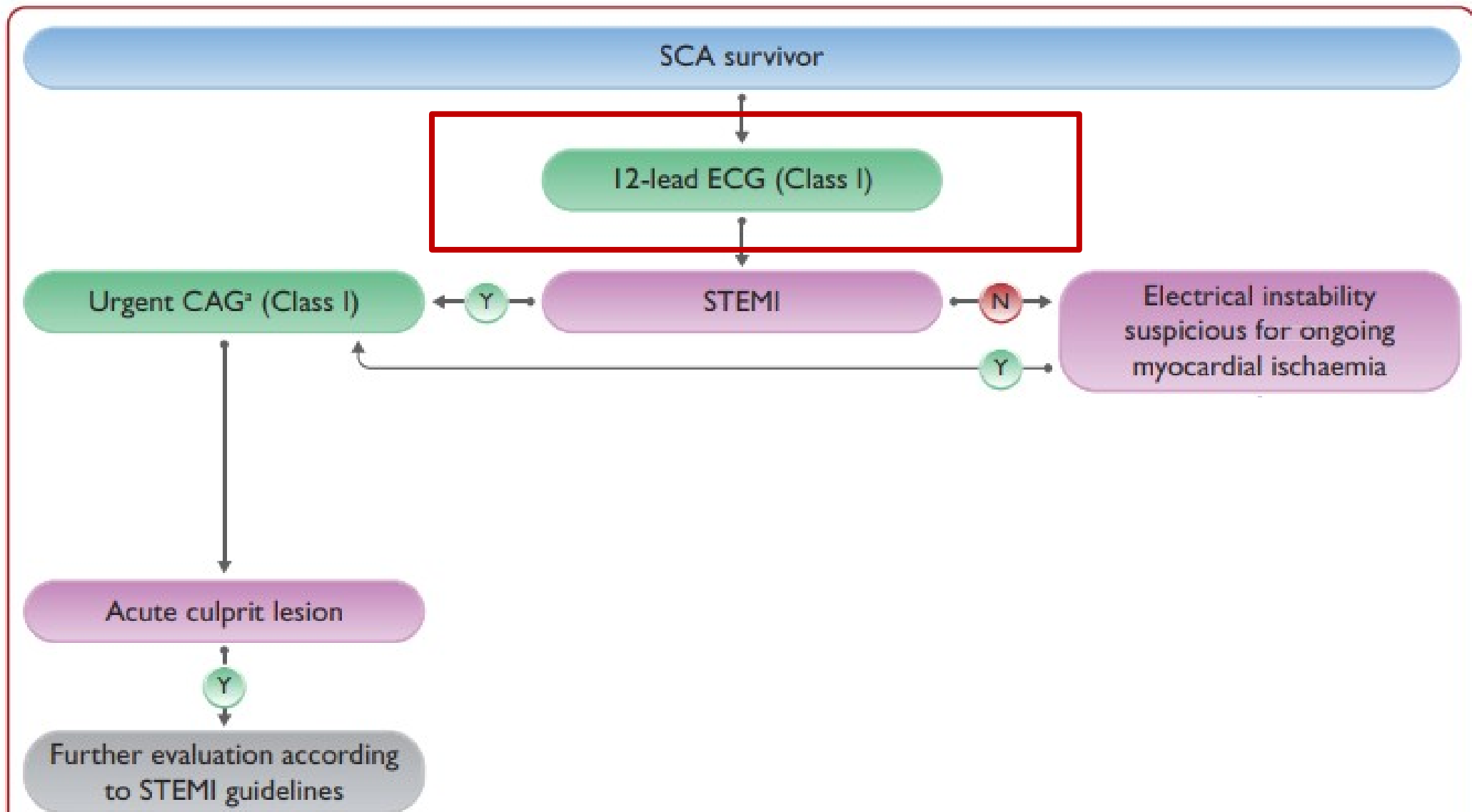


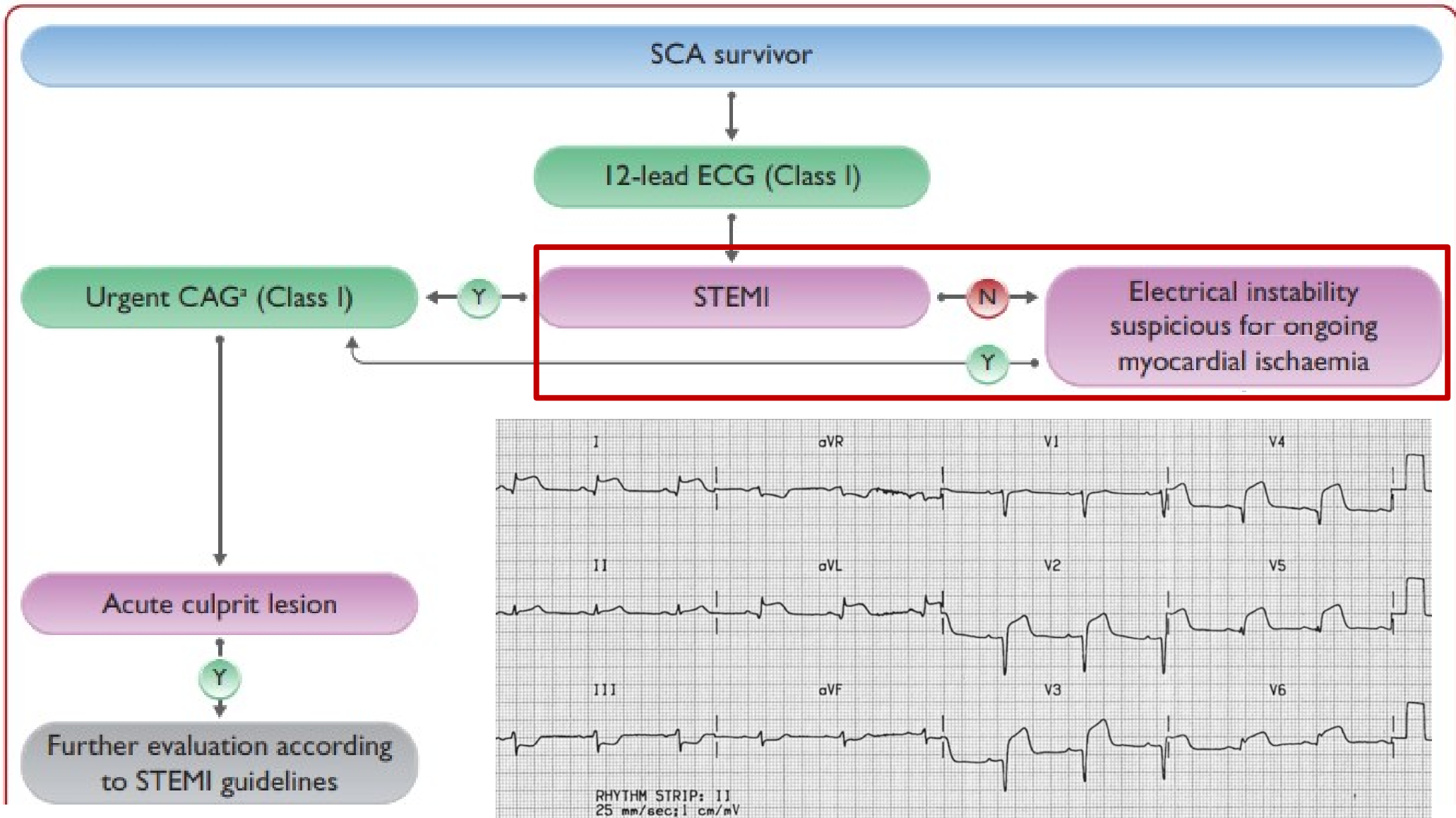
Επιζών από αιφνίδιο καρδιακό  
θάνατο...

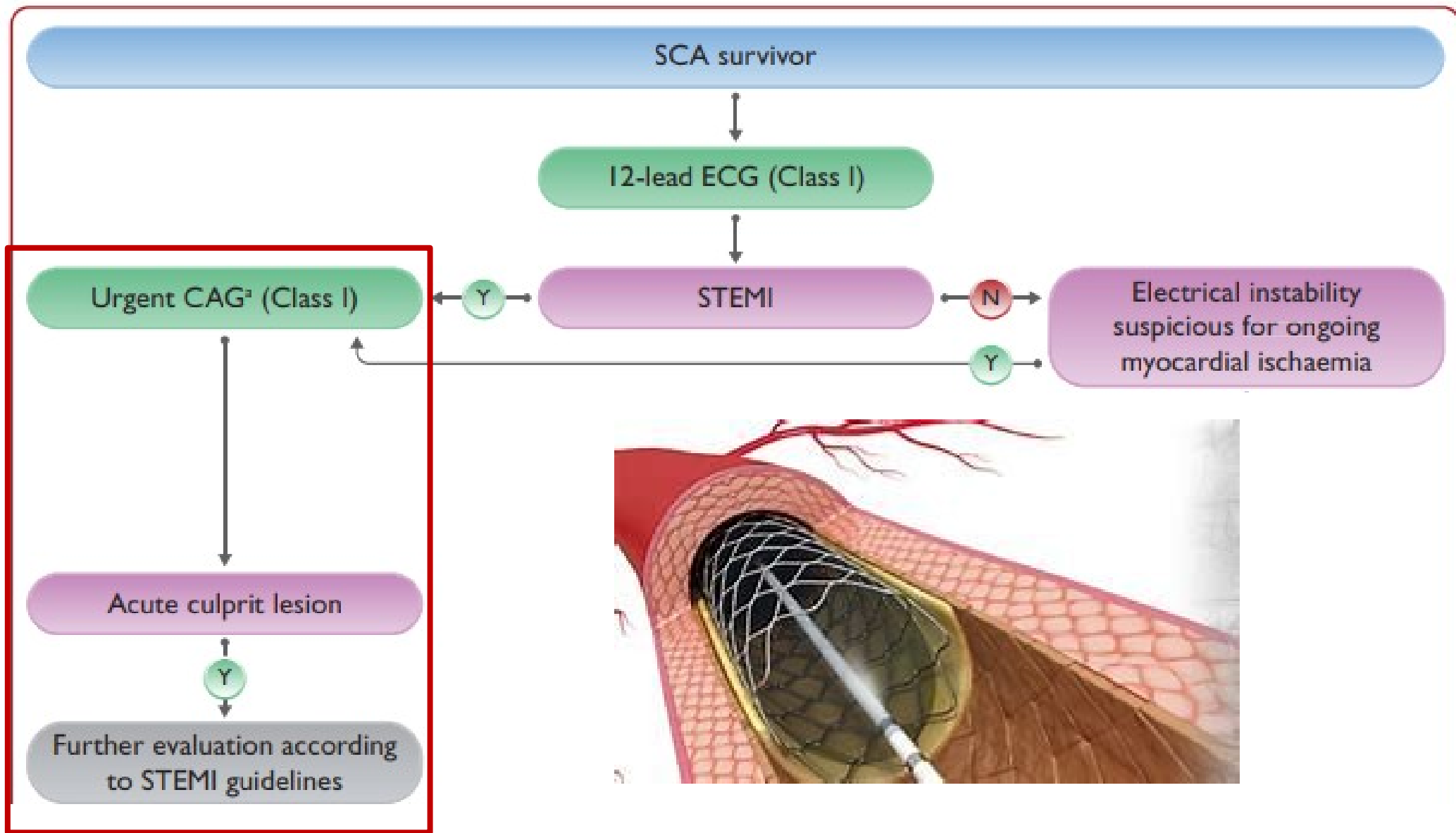
... χωρίς γνωστή στεφανιαία νόσο

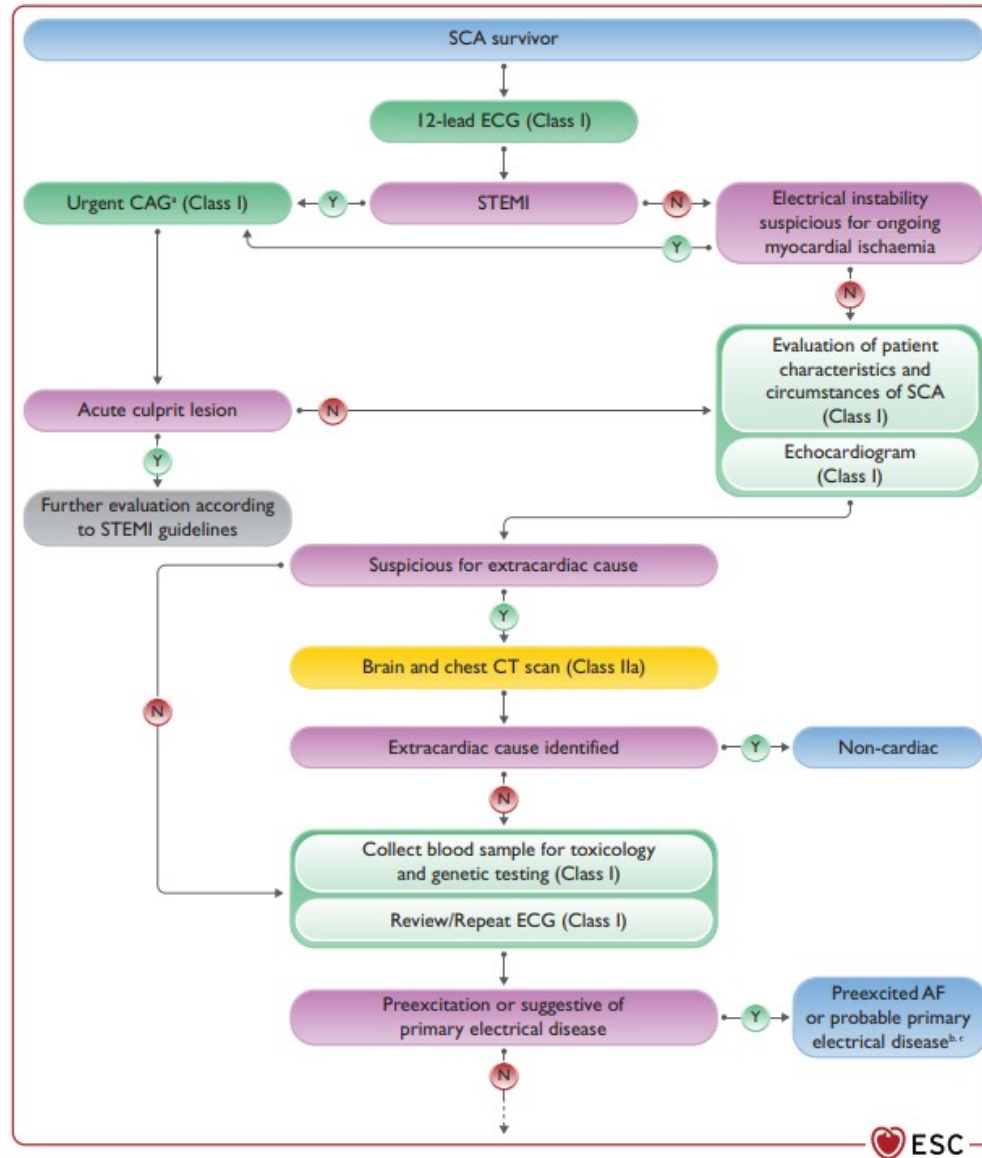


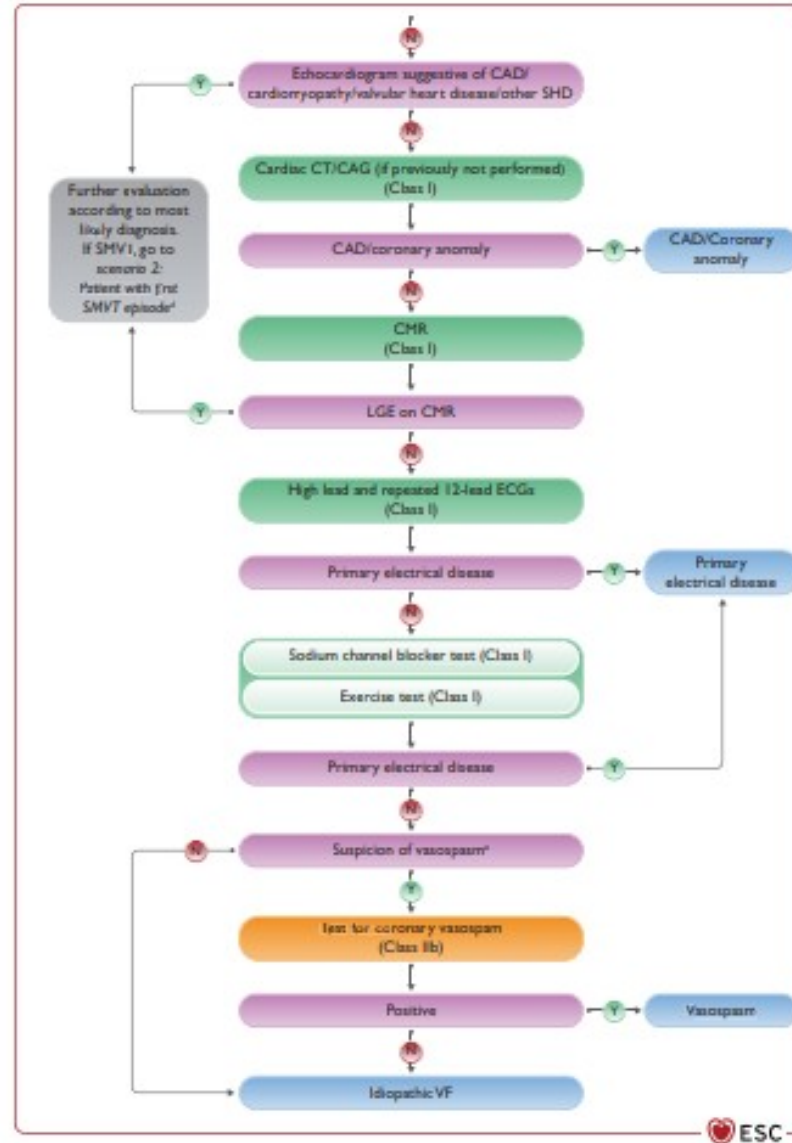


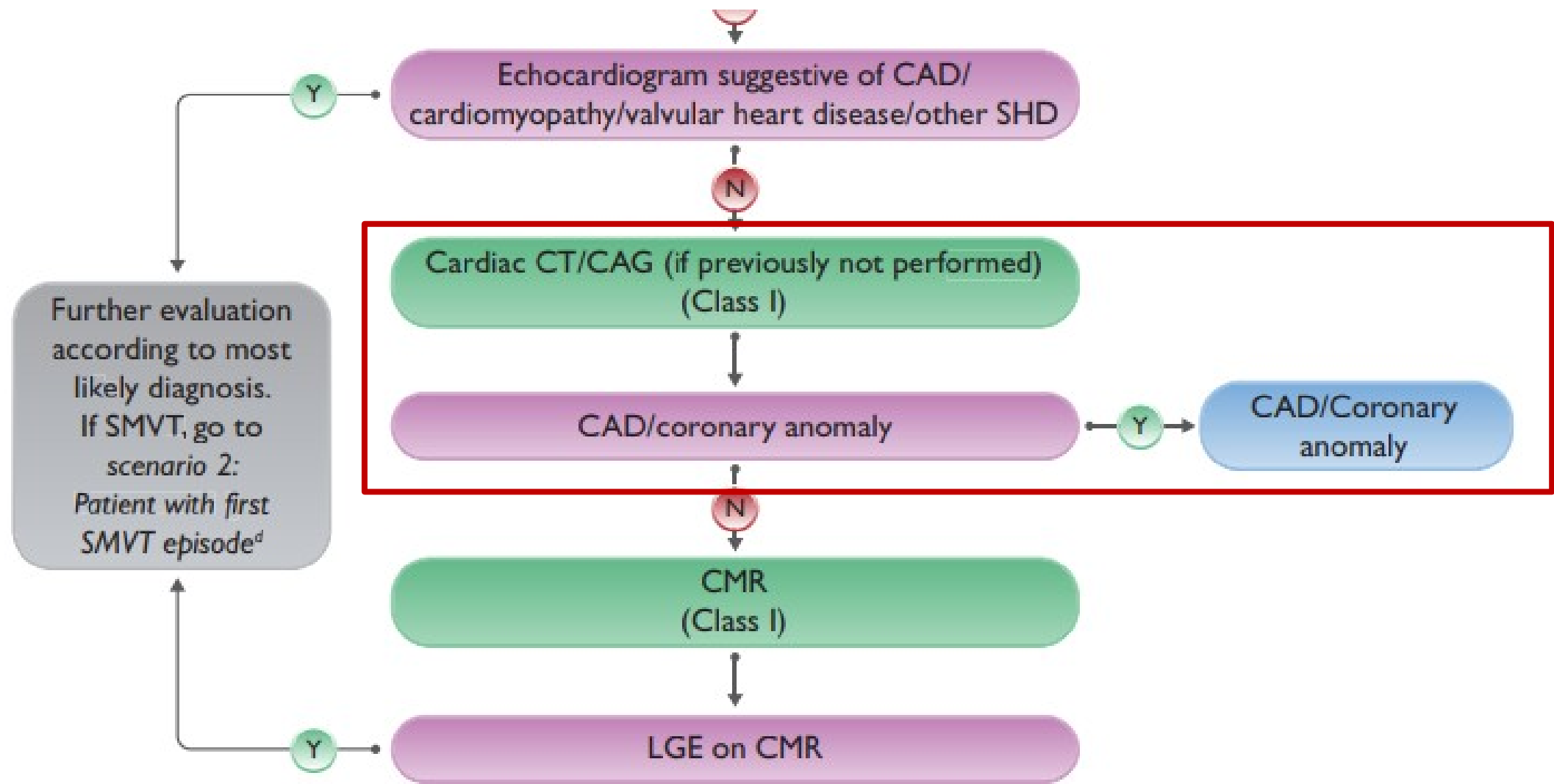








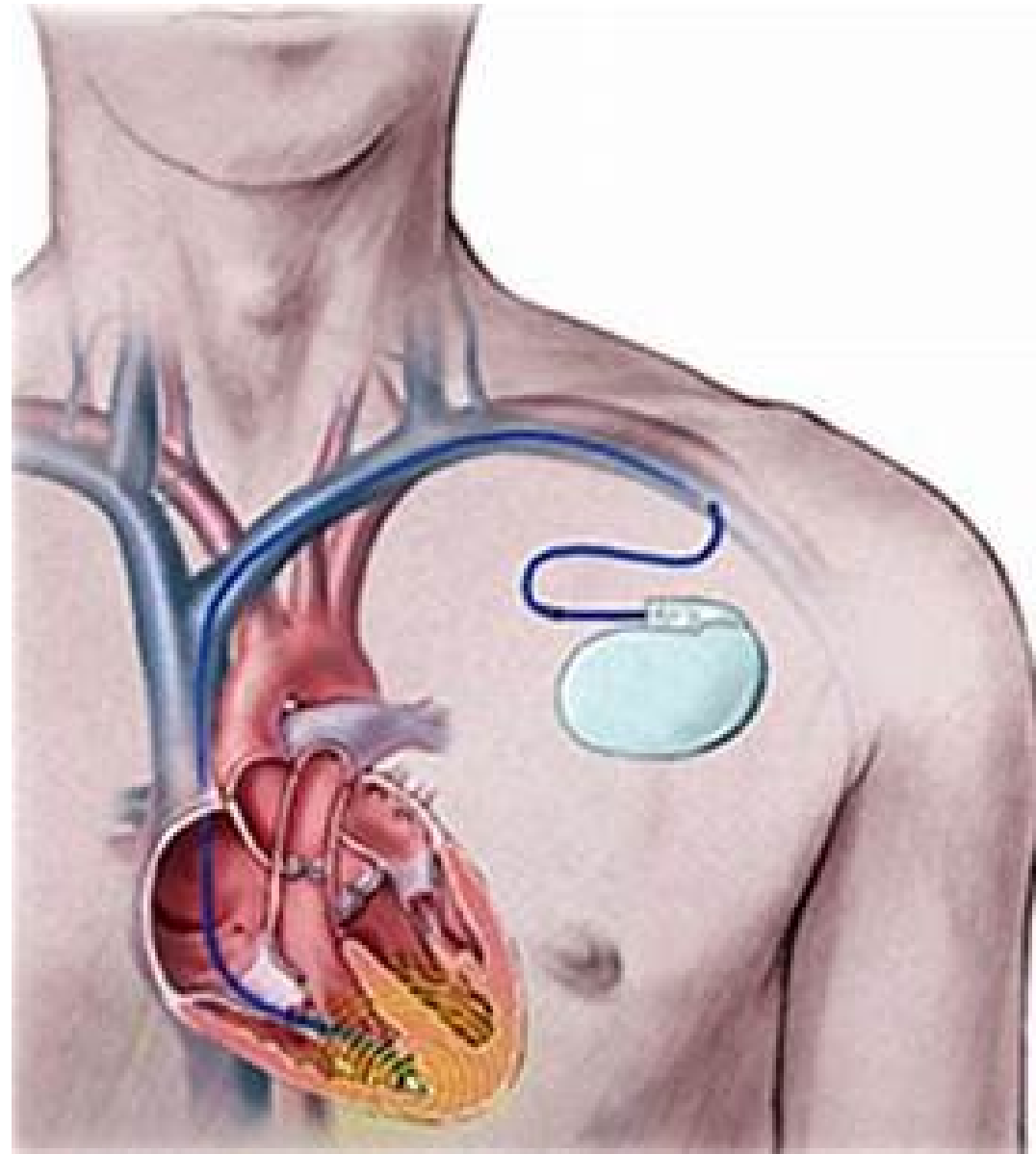




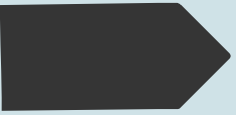


**Prevention**









Σας ευχαριστώ!