

An Update on Heart Failure

Gayathri Kumarasinghe

MBBS BSc(Med) PhD FRACP

ESC Heart Failure guidelines (August 2021)

New terminology

New treatment in HFrEF and HFpEF

Heart failure definitions

HFrEF
Reduced
Ejection Fraction
 $\leq 40\%$

Significant systolic impairment

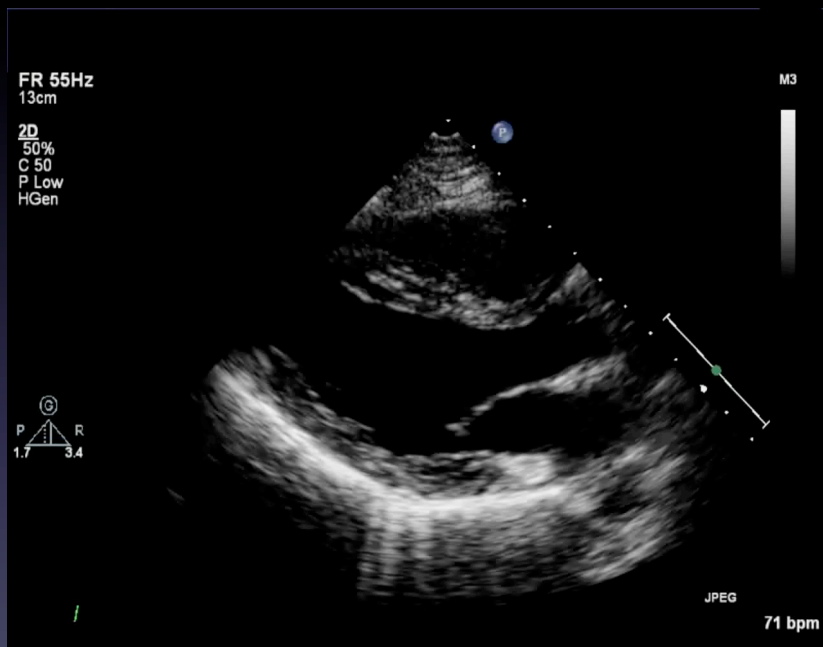
HFpEF
Preserved
Ejection Fraction
 $\geq 50\%$

*Symptoms with structural/functional
cardiac abnormalities and/or raised BNP*

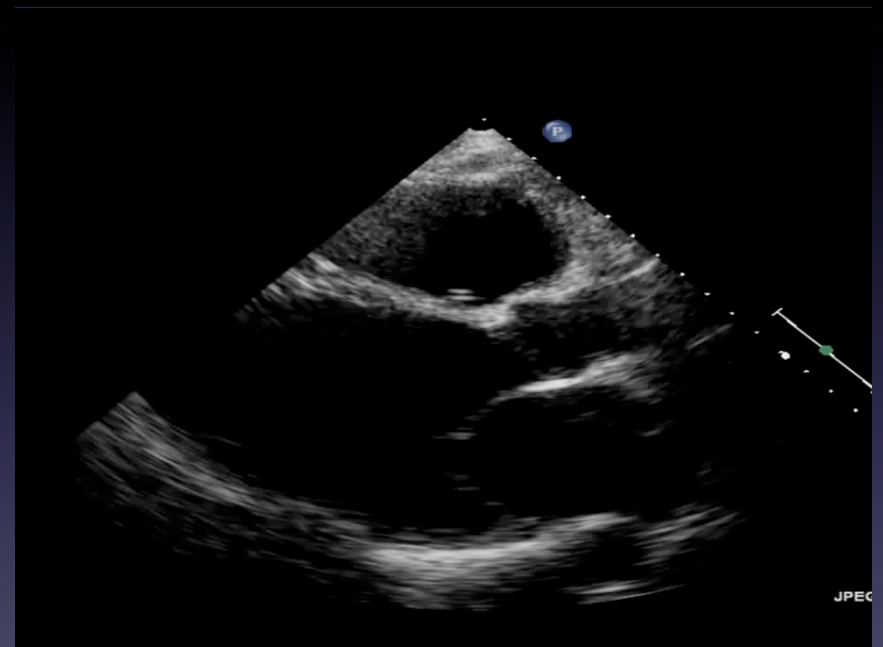
HFmrEF
Mildly Reduced
Ejection Fraction
41-49%

*(previously 'mid-range' EF)
Similar to HFrEF, eg. high ischaemic aetiology, medication response*

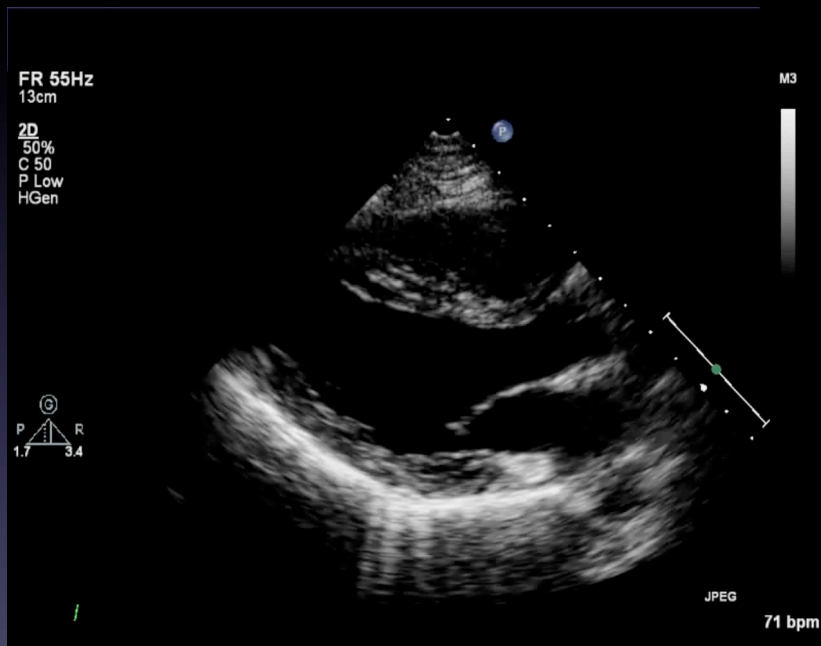
Normal heart



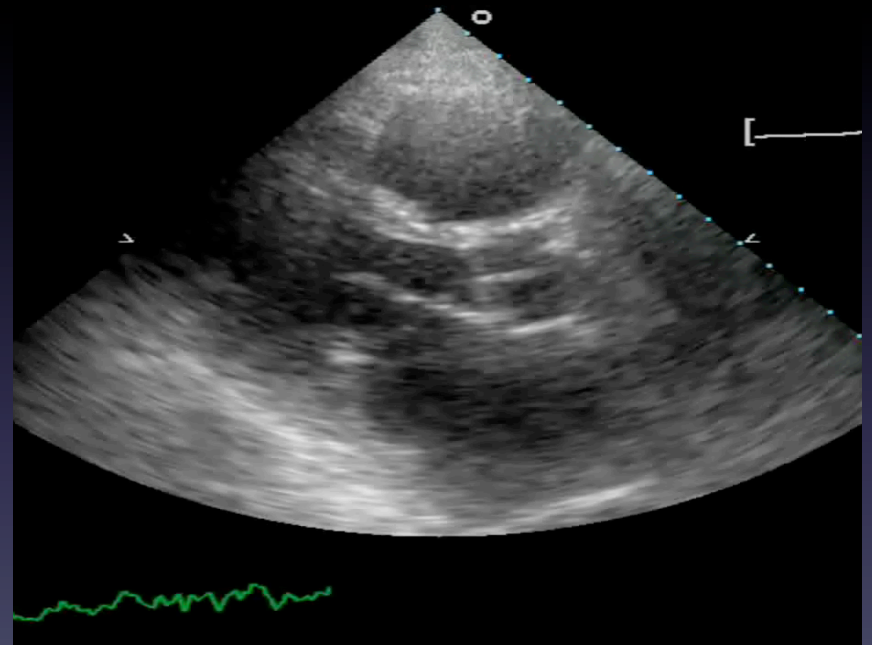
Systolic heart failure (HFrEF)



Normal heart



Diastolic heart failure (HFpEF)



Prognosis in HF

- Mortality still very high
 - 1-year 20% and 5-year 53%
 - Women better survival
 - HFmrEF better survival than HFrEF
 - HFpEF similar survival to HFrEF
- QOL poor
- Average 1 hospitalisation per year
- Higher rate of admissions if diabetes, AF, raised BMI, low eGFR
- Hospitalisation for HF predicted to increase due to aging population

Causes of heart failure

- Coronary artery disease
- Valve disease
- Cardiomyopathies
- Infective
- Infiltrative
- Storage disorders
- Endomyocardial disease
- Pericardial disease
- Hypertension
- Arrhythmias
- Congenital heart disease
- Drug-induced
 - Chemo/immunotherapy
- Metabolic
- Neuromuscular

Causes of heart failure

- Coronary artery disease
- Valve disease
- Cardiomyopathies
- Infective
- Infiltrative
- Storage disorders
- Endomyocardial disease
- Pericardial disease
- Hypertension
- Arrhythmias
- Congenital heart disease
- Drug-induced
 - Chemo/immunotherapy
- Metabolic
- Neuromuscular

Diagnostic tests in suspected heart failure

- BNP/NT-pro BNP
- 12-lead ECG
- Transthoracic echocardiography
- Chest X-ray
- Routine bloods
 - FBC, UEC, TFT, fasting BSL, HbA1c, lipids, iron studies
- Coronary angiography or CTCA, CMR
- Cardiopulmonary exercise testing, right heart catheterisation

Causes of raised BNP or NT-pro BNP

| | |
|--------------------|--|
| Cardiac | Heart failure ACS 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 COPD Severe infections (including pneumonia and sepsis) Severe burns Anaemia Severe metabolic and hormone abnormalities (e.g. thyrotoxicosis, diabetic ketosis) |

BNP < 35 pg/ml

or

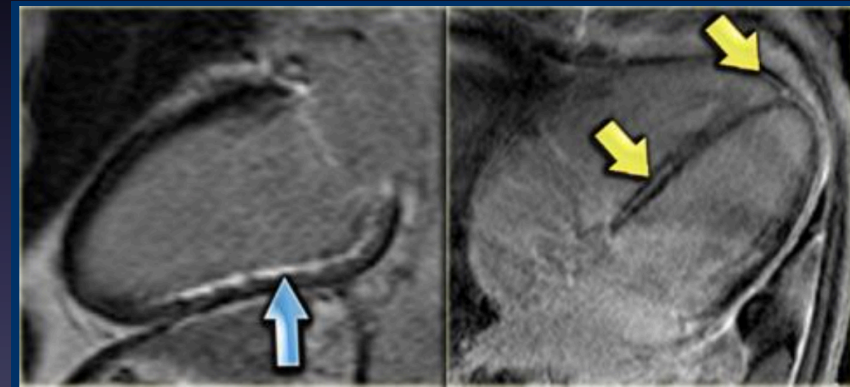
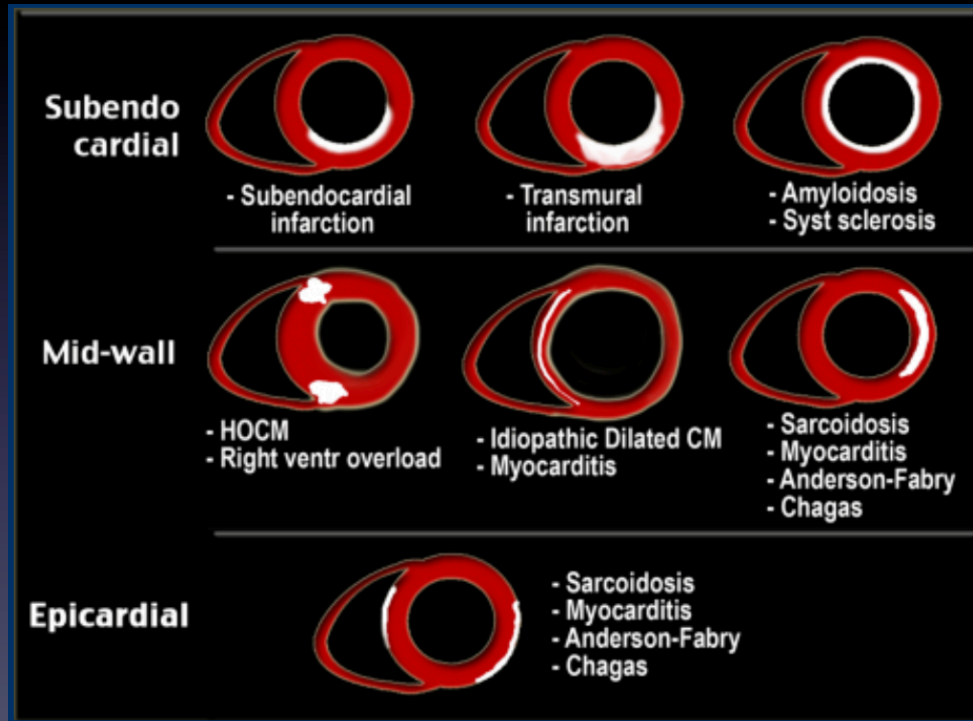
NT-pro BNP < 125 pg/ml

has good negative predictive value

BNP/NT-pro BNP can be very low
in obese patients

Cardiac MRI

Identifying the aetiology of heart failure



Treatment

- Heart failure with reduced EF -

Aims

1. Reduction in mortality
2. Reduction in hospitalisation
3. Improvement in functional capacity, quality of life

HFrEF Pharmacotherapy

Cornerstone therapy

➤ Target RAAS and sympathetic nervous system

ACE-I / ARB or ARNI

Cardioselective beta-blocker

Mineralocorticoid receptor antagonist

Uptitrate to maximum tolerated recommended dose

HFrEF new pharmacotherapy

SGLT2 inhibitors

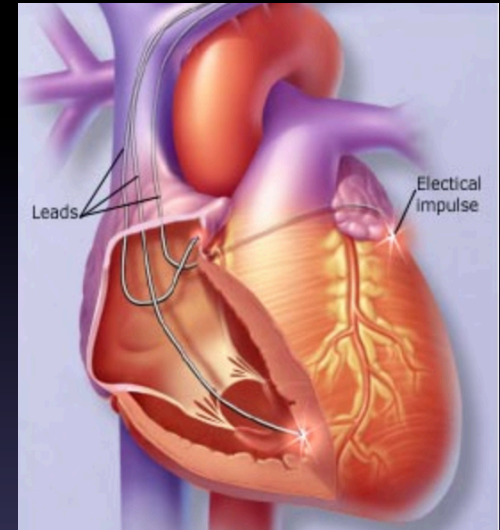
- Dapagliflozin and Empagliflozin
 - Added to ACEI / ARNI / BB / MRA
 - Australian PBS - patients with diabetes only
 - Dapagliflozin – access via PFP
- Reduce the risk of CV death and worsening HF

Improving symptoms of HF

- Diuretics
 - Loop diuretics \pm thiazides
- Ivabradine
 - Sinus rhythm, $HR \geq 70\text{bpm}$, $LVEF \leq 35\%$, hospitalisation within 1 year
- (Digoxin)
 - Digoxin level $< 1.2\text{ng/ml}$

Cardiac resynchronisation therapy

- Symptomatic patients
- $\text{QRS} \geq 150 \text{ ms}$
 - $\text{QRS} \geq 130 \text{ ms}$
- $\text{LVEF} \leq 35\%$ despite optimal medical therapy
- Sinus rhythm or AV nodal ablation in AF



Ventricular arrhythmias in HFrEF

- Amiodarone
 - Reduces ventricular arrhythmias, but no reduction in mortality
- Implantable cardioverter-defibrillators (ICD)
 - Reduces risk of sudden cardiac death
 - Secondary prevention
 - Ischaemic CM > > non-ischaemic CM
 - If LVEF $\leq 35\%$ after 3 months of optimal medical therapy

Treating iron deficiency

Ferritin < 100ug/l

or 100-299 if Trans sat < 20%

Hb 95-135

Ferric carboxymaltose 1g IV

Improves symptoms and reduces HF
hospitalisation

Management of HFrEF

To reduce mortality - for all patients

ACE-I/ARNI

BB

MRA

SGLT2i

To reduce HF hospitalization/mortality - for selected patients

Volume overload

Diuretics

SR with LBBB ≥ 150 ms

CRT-P/D

SR with LBBB 130–149 ms or non LBBB ≥ 150 ms

CRT-P/D

Ischaemic aetiology

ICD

Non-ischaemic aetiology

ICD

Atrial fibrillation

Anticoagulation

Atrial fibrillation

Digoxin

PVI

Coronary artery disease

CABG

Iron deficiency

Ferric carboxymaltose

Aortic stenosis

SAVR/TAVI

Mitral regurgitation

TEE MV Repair

Heart rate SR >70 bpm

Ivabradine

Black Race

Hydralazine/ISDN

ACE-I/ARNI intolerance

ARB

For selected advanced HF patients

Heart transplantation

MCS as BTT/BTC

Long-term MCS as DT

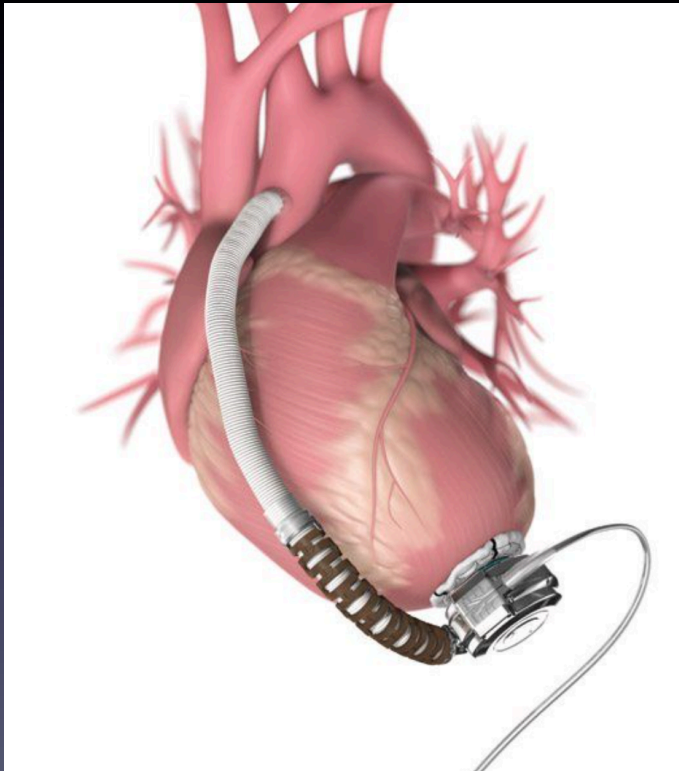
To reduce HF hospitalization and improve QOL - for all patients

Exercise rehabilitation

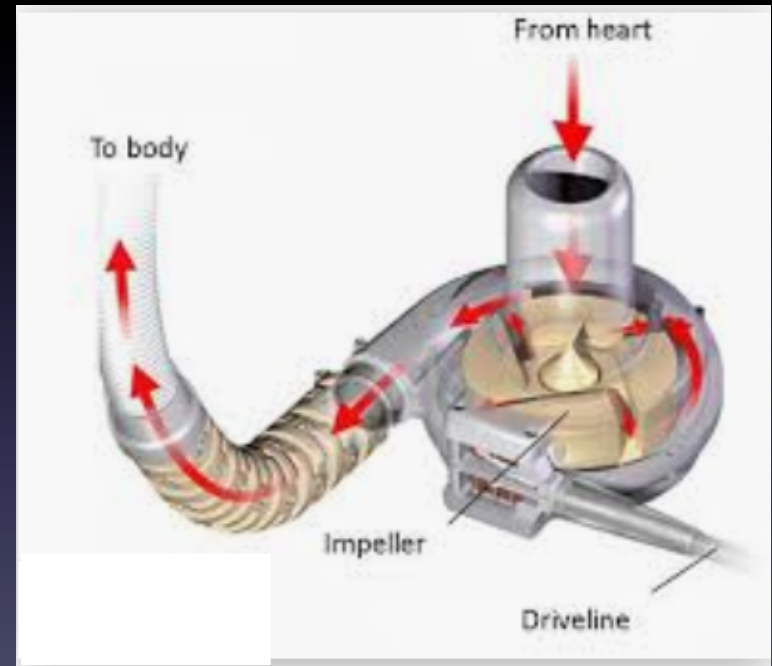
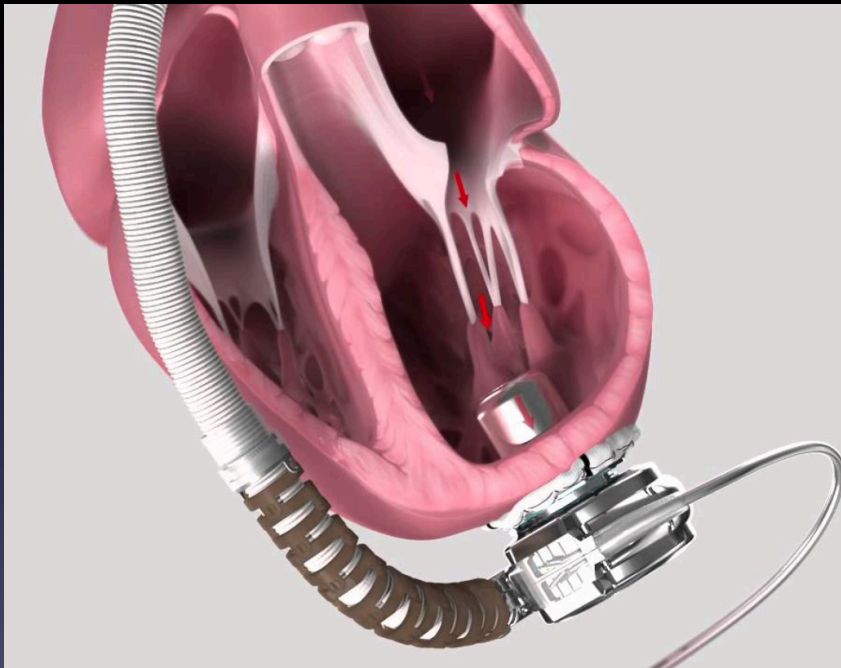
Multi-professional disease management

Ventricular assist devices

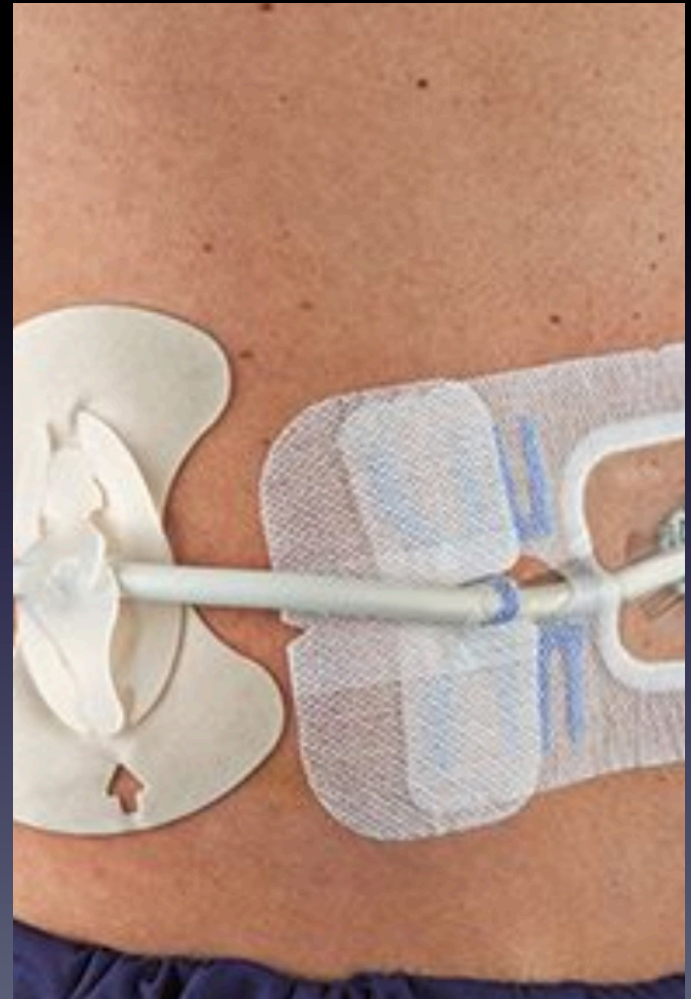
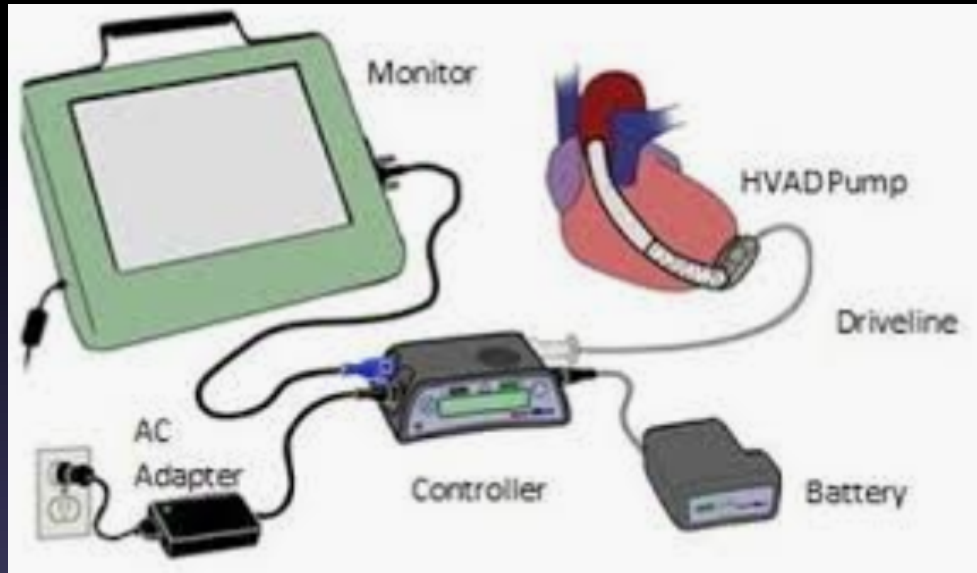
- Left, right or biventricular assist devices



Ventricular assist devices



Ventricular assist devices



Ventricular assist devices

- Indications:
 - Severe intractable heart failure (L, R or biventricular)
 - As a bridge to transplantation
- Complications: GI bleeding, infections, pump thrombosis, haemolysis

Cardiac transplantation

- Patients with advanced heart failure
 - Frequent hospitalisations
 - Symptoms of low cardiac output and congestive heart failure
 - Despite optimal medical and device therapy

HFmrEF

- LVEF 41-49%
- Features of patients similar to HFrEF
 - Men, younger, IHD, less AF and comorbidities
 - Includes patients who improved from LVEF \leq 40% or declined from \geq 50%

HFmrEF

- Diuretics for congestion
- ACE-I, ARB, BB, MRA, ARNI may be considered
 - Often patients on these treatments for other indications, therefore should be continued
- Device therapy – insufficient evidence

HFpEF

- LVEF $\geq 50\%$
- Older patients, female, AF, CKD, non-CV comorbidities more common
- Screen for causes and treat non-CV comorbidities
- Heterogenous condition
- No benefit in ACE-I, ARB, ARNI, BB, MRA
- Diagnostic features:
 - Dilated LA, raised filling pressures ($E/e' > 9$), raised NT-pro BNP, raised pulmonary pressures

HFpEF

SGLT2-I

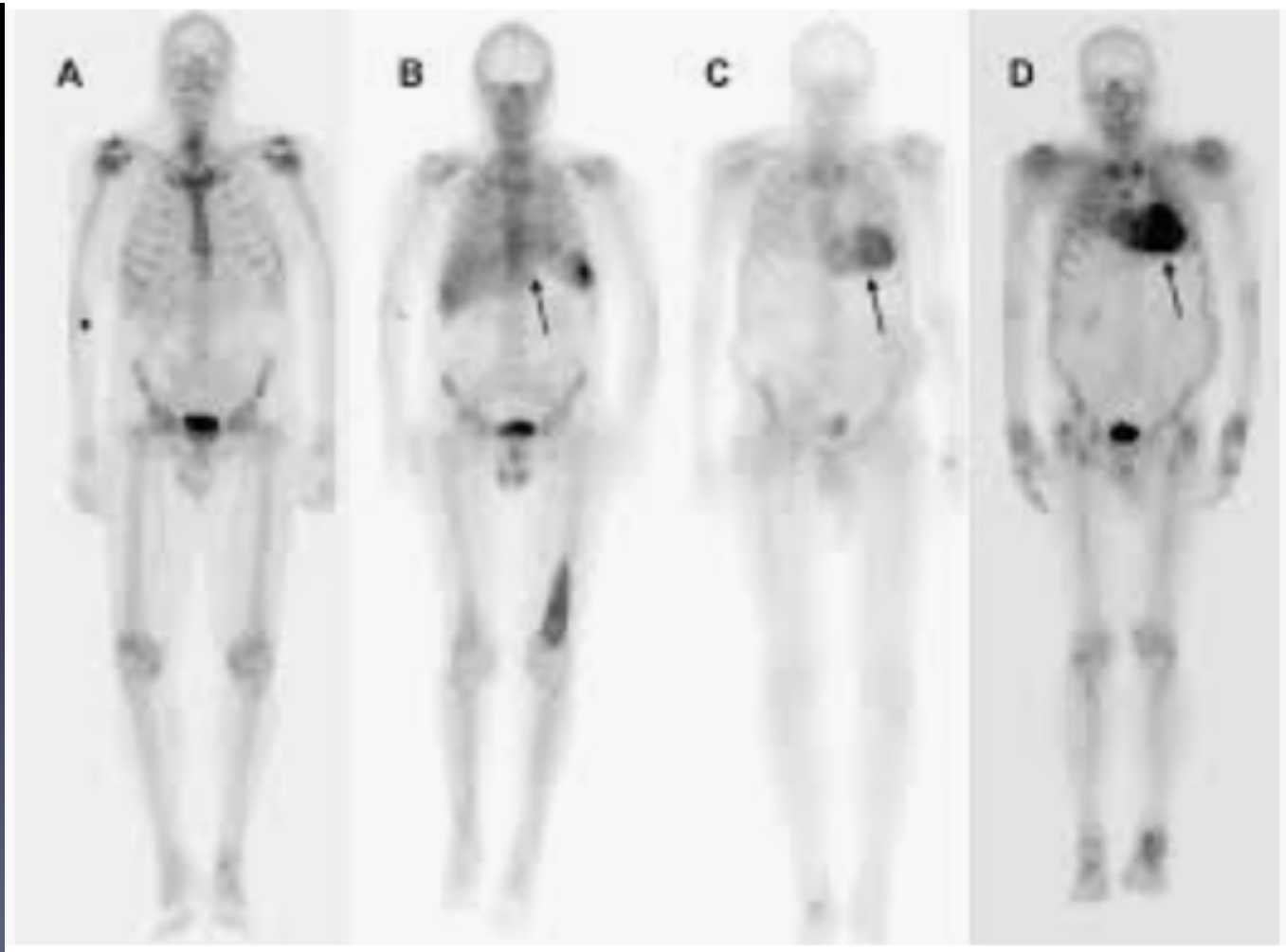
- EMPEROR-Preserved trial
 - Reduced cardiovascular death or hospitalisation
 - However LVEF > 40%
 - Empagliflozin 10mg daily
- The only medication shown to improve survival and hospitalisation in HFpEF.

Cardiac amyloidosis

- Can be a cause of HFpEF
- Suspect if LVH
 - Other clues: peripheral neuropathy, bilateral carpal tunnel, other systemic involvement
 - AL: Serum EPG, IEPG, free light chains, urine Bence Jones protein, bone marrow biopsy
 - ATTR: Bone scan
- If AL amyloid – treat the cause (eg. myeloma)
- If ATTR amyloid
 - Clinical trials underway for RNA interference agents – prevents formation of ATTR protein
 - Tafamidis – Stabilises ATTR tetramer (prevent breakdown into monomer)

Nonbiopsy Diagnosis of Cardiac Transthyretin Amyloidosis.

Gillmore JD¹, Maurer MS¹, Falk RH¹, Merlini G¹, Damy T¹, Dispenzieri A¹, Wechalekar AD¹, Berk JL¹, Quarta CC¹, Grogan M¹, Lachmann HJ¹, Bokhari S¹, Castano A¹, Dorbala S¹, Johnson GB¹, Glaudemans AW¹, Rezk T¹, Fontana M¹, Palladini G¹, Milani P¹, Guidalotti PL¹, Flatman K¹, Lane T¹, Vonberg FW¹, Whelan CJ¹, Moon JC¹, Ruberg FL¹, Miller EJ¹, Hutt DF¹, Hazenberg BP¹, Rapezzi C¹, Hawkins PN¹.



Cardiac amyloidosis

- Can be a cause of HFpEF
- Suspect if LVH
 - Other clues: peripheral neuropathy, bilateral carpal tunnel, other systemic involvement
 - AL: Serum EPG, IEPG, free light chains, urine Bence Jones protein, bone marrow biopsy
 - ATTR: Bone scan
- If AL amyloid – treat the cause (eg. myeloma)
- If ATTR amyloid
 - Clinical trials underway for RNA interference agents – prevents formation of ATTR protein
 - Tafamidis – Stabilises ATTR tetramer (prevent breakdown into monomer)

Thank you