

A patient with acute heart failure and renal impairment

ACCA Masterclass 2017

Dr Sofie Gevaert

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Mister P. J.M., 67-years-old

- **Cardiac risk factors:** Ex-smoker, AHT, Type 2 diabetes, BMI 43, Hyperlipidaemia
- **Medical history:**
 - 2009: Hospitalisation for heart failure: HFpEF
 - 2009: Sick sinus syndrome: DDD pacemaker
 - 1/2016 NSTEMI: PCI D1
 - 2014: Paroxysmal atrial fibrillation
 - COPD GOLD II
 - 11/2011
 - NSTEMI, PCI LAD: 1 month triple R/: VKA-ASA-Clopidogrel
 - Gastro-intestinal bleed with need for transfusion: stop aspirin
 - CKD stage 3B: Serum creatinin 2.09mg/dl, eGFR 30.3mL/min/1.73m²
 - Echocardiography 11/2016:
 - Concentric LVH, EDD 52mm, nl systolic LVF
 - Pseudonormal relaxation pattern, E/E' 16
 - Mild TR: 41mmHg, VCI>17mm, resp variation
 - TAPSE 11mm

12/2016: hospitalization for AHF

- **Complaints:**

Dyspnoea NYHA III, progressive over 6 days after resp infection treated with AB

Peripheral edema

Weight + 5kg

- **Medication:**

- Allopurinol 100mg OD
- Atorvastatin 80mg, OD
- **Bisoprolol 2.5 mg daily**
- **Bumetanide 1mg OD**
- Clopidogrel 75mg
- Pantoprazole 40mg
- **Ramipril 5mg OD**
- Warfarine \approx INR
- Salmeterol/fluticason inhaler

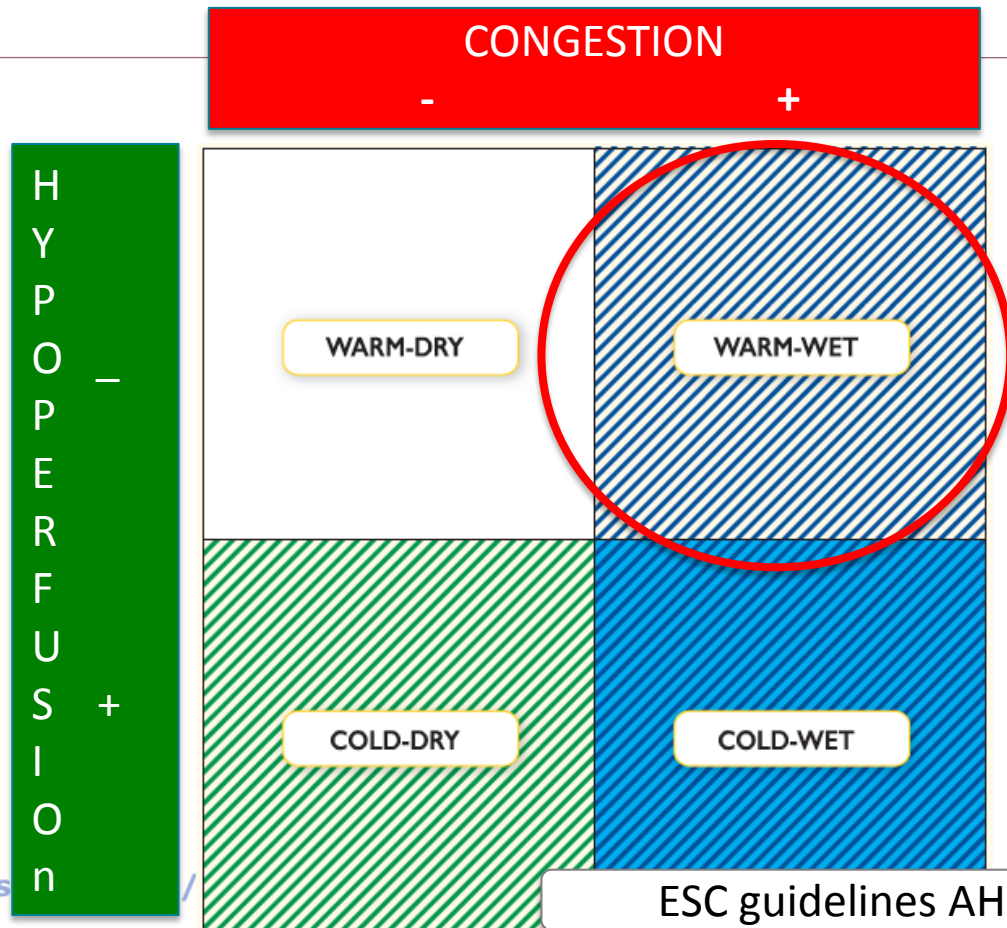
12/2016: hospitalization for AHF

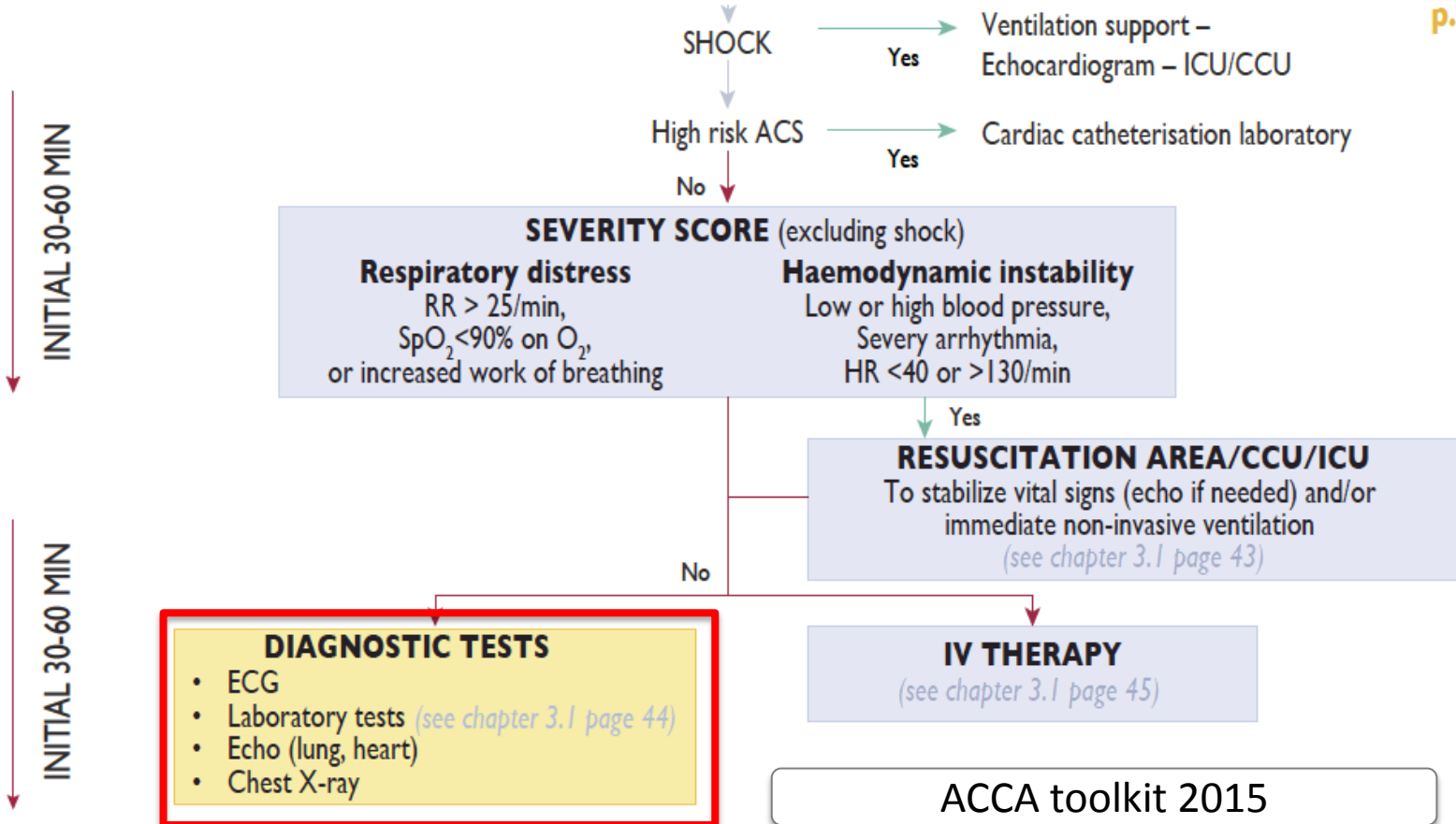
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- **Clinical exam**

- BP 116/60 mmHG, HR 63 BPM, SaO₂ 92%, RR 18
- Jugular distension ++
- Normal heart sounds
- Decreased breathing, rales
- Peripheral edema ++

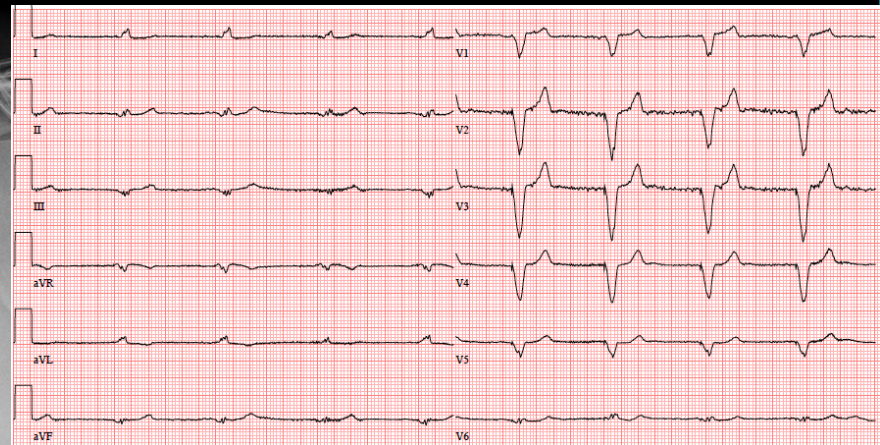
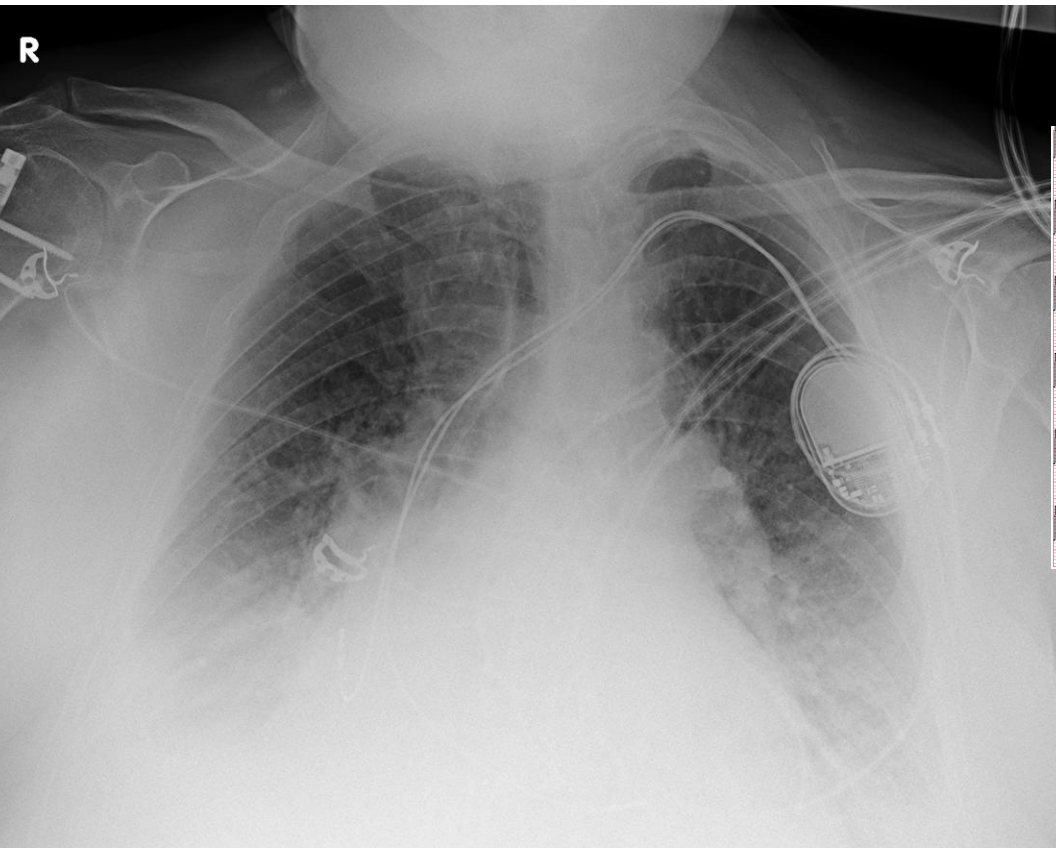
Clinical Profile?





Chest X-ray-ECG

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- **Lab results**

- Hct 36.3% (39-49)
- K⁺ 5.5 mmol/L
- Serum Creatinin **2.85 mg/dL** (0.72-1.17), eGFR: 22 mL/min/1.73 m²
- proBNP: 2,400 pg/mL (≤125)
- PH 7.37, pO₂ **55**, pCO₂ 33.4

Timing echocardiography?

- Immediately
- <48 hours of admission
- Before discharge
- Not necessary, receive

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). Pocket-size echocardiography may be used as an extension of the clinical examination in the first instance where available. Repeated echocardiography is usually not needed unless there is relevant deterioration in clinical status.

Echocardiography

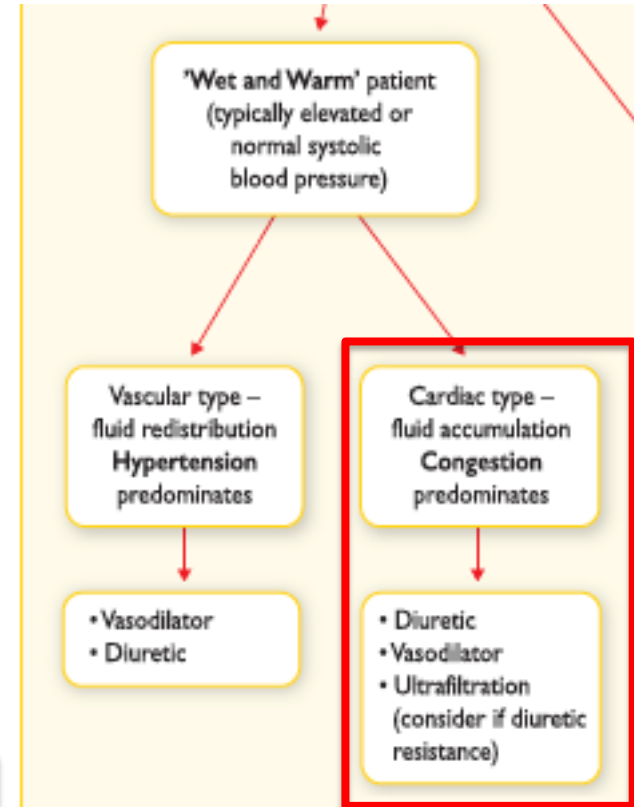
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- Moderate views
- Restrictive relaxation pattern, E/E' med 26
- TR grade 2+: 54mmHg, VCI>17mm no resp. variation
- TAPSE 11mm

→ ADHF triggered by respiratory infection

Initial treatment?

- IV diuretics
- IV diuretics and vasodilators
- Ultrafiltration



Diuretics-Ultrafiltration

Diuretics

Intravenous loop diuretics are recommended for all patients with AHF admitted with signs/symptoms of fluid overload to improve symptoms. It is recommended to regularly monitor symptoms, urine output, renal function and electrolytes during use of i.v. diuretics.

I

C

In patients with new-onset AHF or those with chronic, decompensated HF not receiving oral diuretics the initial recommended dose should be 20–40 mg i.v. furosemide (or equivalent); for those on chronic diuretic therapy, initial i.v. dose should be at least equivalent to oral dose.

I

B

It is recommended to give diuretics either as intermittent boluses or as a continuous infusion, and the dose and duration should be adjusted according to patients' symptoms and clinical status.

I

B

Combination of loop diuretic with either thiazide-type diuretic or spironolactone may be considered in patients with resistant oedema or insufficient symptomatic response.

IIb

C

Ultrafiltration may be considered for patients with refractory congestion, who failed to respond to diuretic-based strategies.

IIb

B

Renal replacement therapy should be considered in patients with refractory volume overload and acute kidney injury.

IIa

C

Referral CCU: IV diuretics and vasodilators

- Continuous infusion bumetanide 0.5mg/h for 24h
- Continuous infusion isosorbide dinitrate 2mg/h
- Negative fluid balance -3L 1st 24h → oral diuretics at increased dose

→

- Improvement kidney function: **2.0 mg/dl (eGFR 34mL/min)** at discharge
- Discharge after 72h:
 - Weight -5 kg
 - Oral dose Bumetanide increased to 2mg
 - Daily weight, salt restriction
 - Follow-up 4 weeks

Readmission 1 week later

- **Complaints:**

Dyspnoea NYHA III, progressive 3 days, peripheral edema, weight +4 kilo

- **Medication:**

- Allopurinol 100mg OD
- Atorvastatine 80mg, OD
- Bisoprolol 2.5 mg daily
- **Bumetanide 2mg OD**
- Clopidogrel 75mg
- Pantoprazole 40mg
- Ramipril 5mg OD
- Warfarine \approx INR

- **Clinical exam**

- BP 128/55 mmHG, HR 56 BPM, SaO₂ 94%
- Jugular distension
- Normal heart sounds
- Rales basal
- Moderate peripheral edema

- **Lab results**

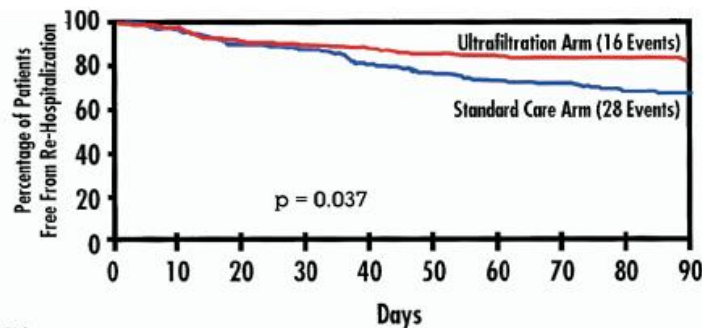
- Serum creatinin:2.3 mg/dL (+0.3mg/dL, GFR 28mL/min)

Could UF have prevented rehospitalization?

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ADHF

Adjustable UF (up to 500mL/h)
VS.
diuretic R/ at discretion physician



| No. Patients at Risk | 0 | 10 | 20 | 30 | 40 | 50 | 60 | 70 | 80 | 90 |
|----------------------|----|----|----|----|----|----|----|----|----|----|
| Ultrafiltration Arm | 88 | 85 | 80 | 77 | 75 | 72 | 70 | 66 | 64 | 45 |
| Standard Care Arm | 86 | 83 | 77 | 74 | 66 | 63 | 59 | 58 | 52 | 41 |

UNLOAD trial,
Costanzo et al, JACC 2007

Compliance problem? Diuretic resistance? Decongestion treatment?

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- **IV diuretics**
- **Increase oral dose**
- **Add thiazide or other**
- **Ultrafiltration**

Referral to ward for IV diuretics

| day | Bumetanide | Diuresis (mL) | Serum creatinine |
|-------|------------------|-----------------|------------------|
| 1 | 20.00h: 0.48mg/h | 1.950 | 2.30 |
| 2 | 0.48mg/h IV | 2.700 | 2.20 |
| 3 SAT | Stop IV 23.00h | 260 (complete?) | 2.51 |

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| 4 SUN | 1 mg PO | 700 | - |
| 5 | 1mg PO | 610 | 3.91 |

Referral to ward for IV diuretics

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| 2 | 0.48mg/h IV | 2.700 | 2.20 |
| 3 SAT | Stop IV 23.00h | 260 (complete?) | 2.51 |
| 4 SUN | 1 mg PO | 700 | - |
| 5 | - | 610 | 3.91 |
| 6 | - | 1.150 | 3.4 |
| 7 | - | 1.500 | 2.45 |
| 8 | - | 1.450 | 1.89 |
| 9 | 1mg PO | 1.500 | 1.83 |

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Too rapid decongestion → AKI

KDIGO definition of AKI

| | |
|---|------|
| Increase in Scr level of ≥ 0.3 mg/dL (26.5 $\mu\text{mol/L}$) | <48h |
| Increase in Scr level of ≥ 1.5 times baseline | <7 d |
| Urine output $<0.5\text{mL/Kg/h}$ | >6h |

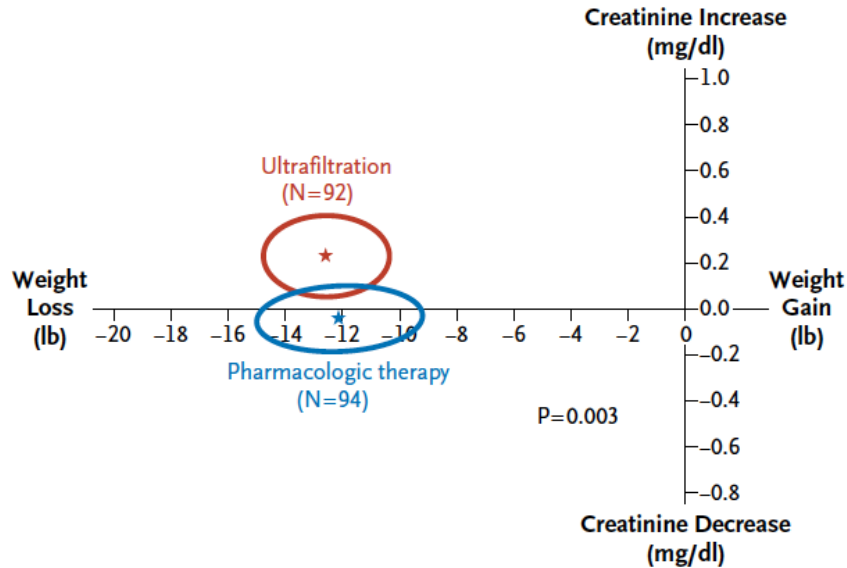
KDIGO staging of AKI

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| Stage | Scr increase | Urine output |
|-------|--|---|
| 1 | $\geq 1.5-1.9$ times baseline or ≥ 0.3 mg/dL | < 0.5 mL/kg/h for $\geq 6-12$ h |
| 2 | ≥ 3 times baseline or ≥ 4 mg/dL or RRT | < 0.5 mL/kg/h for ≥ 12 h |
| 3 | ≥ 3 times baseline or ≥ 4 mg/dL or RRT | < 0.3 mL/kg/h for ≥ 24 h or Anuria for ≥ 12 h |

CRS type 1: acute cardiorenal

Could UF have prevented AKI ?



ADHF+WRF:
Fixed UF rate (200mL/h)
vs.
defined stepped pharmacological R/

Figure 1. Changes in Serum Creatinine and Weight at 96 Hours (Bivariate Response).

CARESS trial,
Bart et al, NEJM 2012

ss 2017

OBJECTIVES The AVOID-HF (Aquapheresis versus Intravenous Diuretics and Hospitalization for Heart Failure) trial tested the hypothesis that patients hospitalized for HF treated with aquapheresis would have a shorter time to first HF event within 90 days after hospital discharge than those treated with intravenous diuretics.

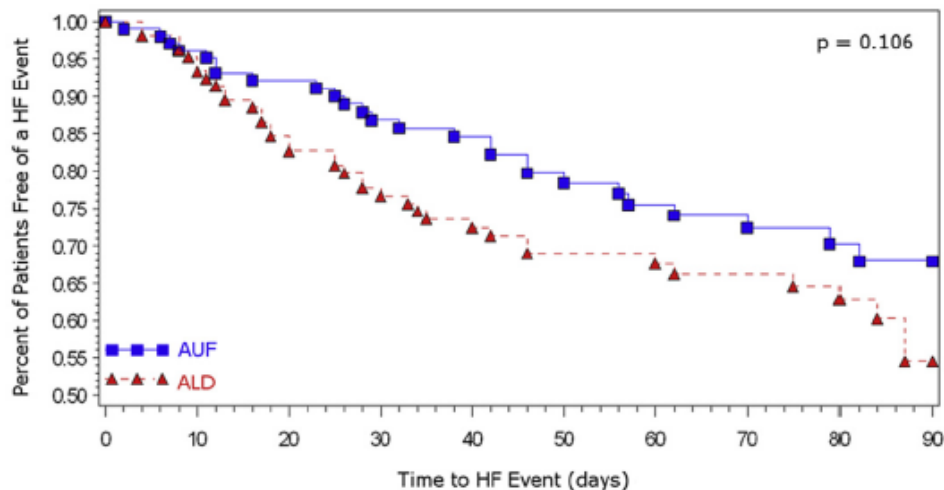
BACKGROUND Congestion in hospitalized heart failure (HF) is associated with increased mortality and morbidity.

METHODS The AVOID-HF trial, designed as a multicenter, randomized controlled trial, was terminated unilaterally and prematurely by the sponsor (Baxter) because of a high rate of adverse events in patients (27.5%). Aquadex FlexFlow System (Baxter HealthCare Corporation) was used for aquapheresis in the randomized treatment, adjudicated whether 90-day mortality was similar.

RESULTS A total of 110 patients were randomized to aquapheresis (AUF) or intravenous diuretics (ALD). The median time to first HF event for the AUF and ALD group were 45 and 35 days, respectively. In patients treated with the ALD group, the AUF group had fewer HF and cardiovascular events. More patients in the ALD group experienced an adverse effect of special interest (p = 0.026). The 90-day mortality was similar.

CONCLUSIONS Compared with the ALD group, the AUF group had a shorter time to first HF event within 90 days and fewer HF and cardiovascular events. More patients in the ALD group experienced a product-related adverse event. Due to the trial's untimely termination, the results are inconclusive. (J Am Coll Cardiol HF 2016;4:95-105) © 2016 by the American College of Cardiology

FIGURE 2 Primary Endpoint: Time to Heart Failure Event after Discharge



| | Baseline | 30 Days | 60 Days | 90 Days |
|-----|----------|---------|---------|---------|
| AUF | 105 | 80 | 52 | 19 |
| ALD | 108 | 74 | 49 | 15 |

No difference in changes kidney function

AVOID trial,
Costanzo, Bart et al, JACC HF 2016

3 weeks later, third admission

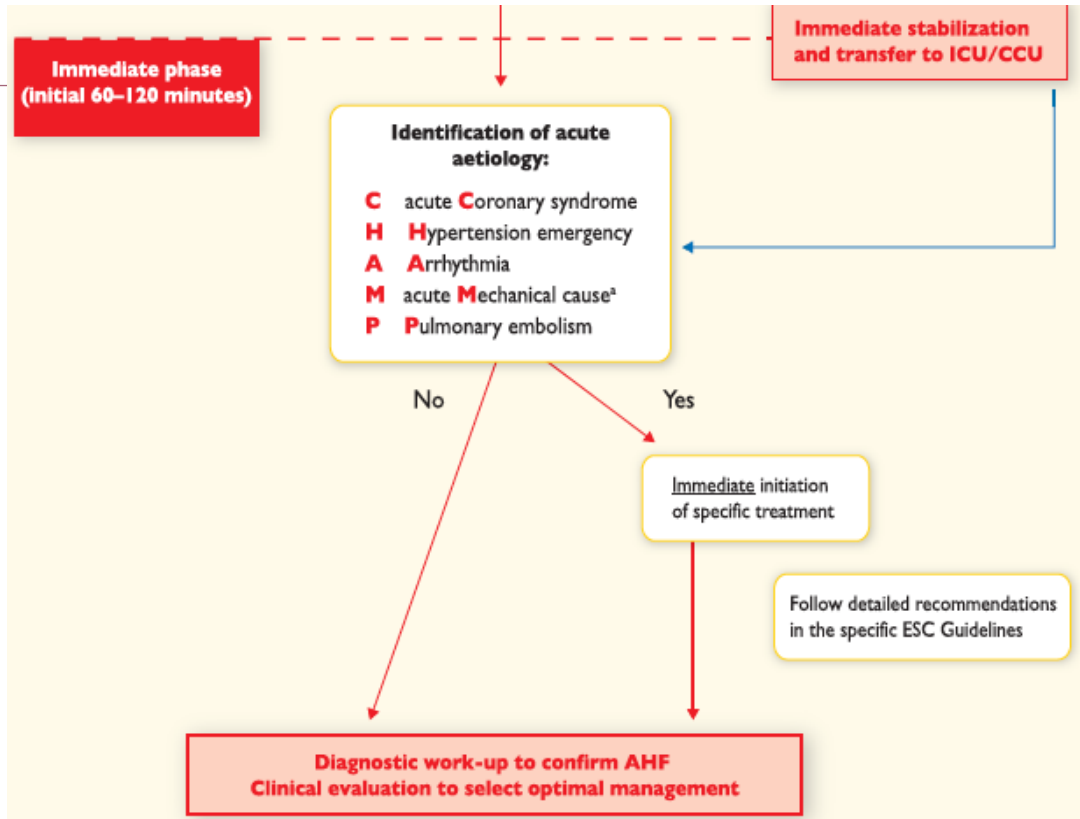
- **Complaints:**

Dyspnoea NYHA IV, progressive since discharge, peripheral edema, weight +4 kg

- **Medication:**

- Allopurinol 100mg OD
- Amlodipine 5 mg OD
- Atorvastatine 80mg, OD
- Bisoprolol 2.5 mg daily
- **Bumetanide 2-3mg OD, depending on weight**
- Clopidogrel 75mg
- Pantoprazole 40mg
- Ramipril 5mg OD
- Warfarine \approx INR

- **Clinical exam**
 - BP 116/60 mmHG, HR 63 BPM, SaO₂ 88%
 - Jugular distension
 - New systolic murmur?
 - Decreased breathing, rales
 - Peripheral edema
- **Lab results**
 - Serum creatinin 2.2mg/dL



UZG Hartbewaking

TIS 0.1 14:54:44

TEE RETRO
X7-2t
46 Hz
8.0cm

Pat. T: 37.0 °C
TEE T: 39.5 °C

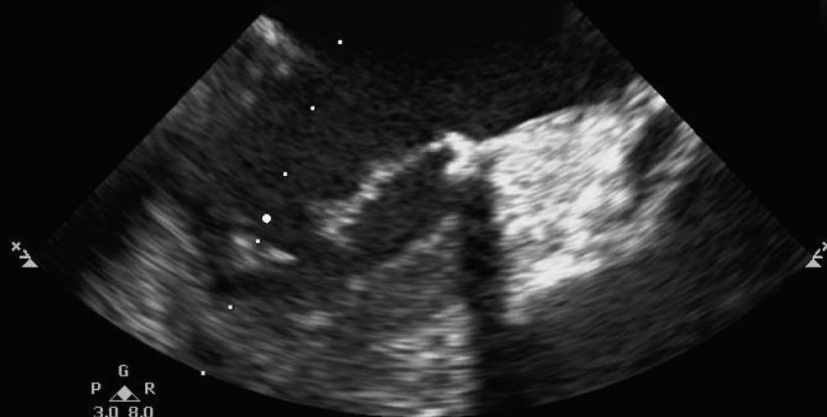
0 83 180



P

2D

Gen
Gn 50
C 48
4/4/0
50 mm/s



G
P R
3.0 8.0

UZG Hartbewaking

TIS 0.8

RO

Pat. T: 37.0 °C

TEE T: 39.9 °C

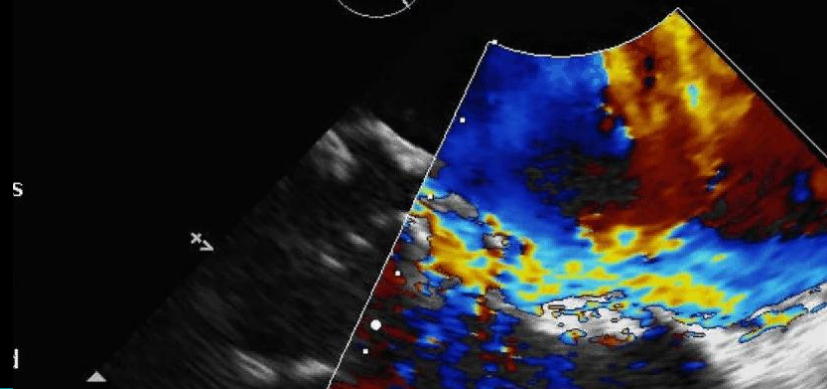
0 127 180



P

S

I



RUPTURED CHORDA of ant. LEAFLET MV

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- Cardiac surgery
- Repair not succesful → bioprosthesis CE Magna Ease
- Postoperative:

AKI, anuria → Dialysis

Thank you

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How to avoid too rapid/much decongestion?

Decongestion goal CLOSE monitoring of ?

- BP
- weight
- Δ renal function/urine output
- Monitoring of intravascular volume
 - Hct as surrogate marker?
- Monitoring of CO?
 - CVO_2 as surrogate marker of CO?
- ...?

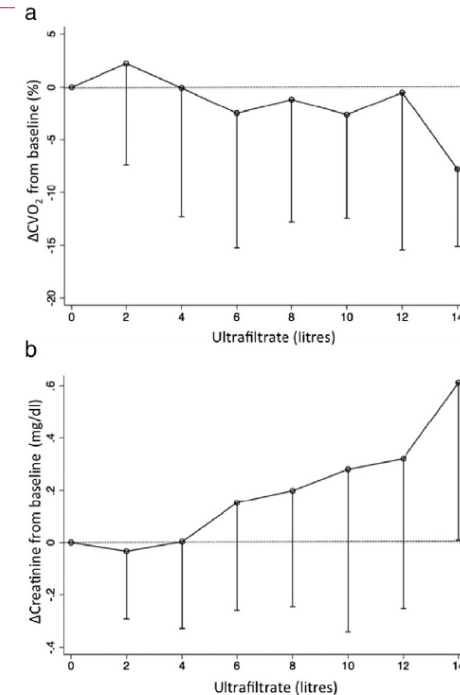


Fig. 1. Title: Mean and standard deviation of a) changes in central venous oxygen level (ΔCVO_2) and b) changes in creatinine (Δ creatinine) from baseline values for every 2 l of ultrafiltrate removed. This figure demonstrates that as the initial 2 l of fluid was removed, there was a rise in mean ΔCVO_2 corresponding to a fall in mean Δ creatinine. Subsequently with further fluid removal beyond 4 l, there was a fall in mean ΔCVO_2 which corresponded to a rise in mean Δ creatinine.