

PROGRAM

9th edition **Live and online Forum Hybrid from Nice, France**

12th & 13th Oct. 19th Nov. 2021

3 days of interactive discussions

Understanding the heart failure guidelines development process and its implications

Live location: Nice, France

Why a Guidelines Forum on heart failure?

9th edition Live & online Forum



The quality of research and development efforts made by manufacturers of pharmacological agents, biomarkers, and medical devices has obviously had a major impact on the position of these products in clinical guidelines on the management of heart failure. Conversely,

the success and failure stories that inevitably accompany the release of new guidelines may be a source of inspiration for further research and development. And, ultimately, guidelines really shape clinical practice if doctors and their patients adopt them.

A better understanding of the processes involved in guidelines development will help companies that develop drugs, biomarkers, and devices to adapt and improve their research and development programmes for the future.

This could also translate into earlier and more efficient development of diagnostic and therapeutic tools for patients.

How do guideliners deal with the available evidence and what are their expectations in this respect? What criteria must a product or an intervention meet to be recommended by heart failure guidelines? What are the most frequent reasons for failure? Why may guidelines of geographically distinct origins but dealing with the same evidence occasionally reach different

Format

A series of short presentations and discussions with a focus on practical examples, case studies and interactivity between Faculty members.

Who should attend?

Attendees include industry representatives with interest in clinical trials, product development, market access, guidelines development and implementation, patients' advocacy specialists, clinical research organizations and payers.

About TMA

The Translational Medicine
Academy (TMA) is an
international academic
Foundation with focused
activities in Mission Critical Areas
of high public health importance
having significant professional
and patient therapeutic and
educational gaps.

Why a Heart Failure Guidelines Forum?

The objective is to better understand the existing Heart Failure guidelines development process and its implications in clinical practice. It is not to develop new guidelines, but to gather a multidisciplinary panel of leading international experts to discuss the latest evidence, ongoing research and controversial issues that have implications for the development of clinical practice guidelines.

Educational objectives

At the end of the meeting, all participants will have a better understanding of the unmet needs and knowledge gaps that need to be addressed by future research and development efforts, and of the most appropriate ways of progressing to conciliate and take advantage of clinical, societal, evidence-based medicine and regulatory requirements.

conclusions? What should be done to avoid that guidelines being lost in translation? How can we adapt the guidelines for low and middle income countries?

Speakers and discussants who will address these topics have been selected from among the top-level North American and international experts in cardiology who participate in the regular update of guidelines. The faculty also comprises clinicians with a strong interest in translational and patient-centered medicine.

The Heart Failure Guidelines Close Forum will be strictly academic – meaning that attendees will not participate during the main sessions and discussions. However, they will be able to transmit written questions to the faculty.

Hybrid forum

A live meeting is scheduled to be held in Nice, France according to regulations.

Remote access will be possible via a secure digital and interactive plateform.

This event is registered with MedTech Europe and declared (provisionally) compliant: https://www.ethicalmedtech.eu/

Program





OCTOBER 12th, 2021 - 14:30-19:30 CEST

14:30–14:35: Welcome / Introductions S. Anker

14:35–15:10: Session 1 – Heart failure guideline process – Rules of engagement

Chairs: S Anker / G Rosano / P Seferovic

• How are the Europe guidelines developed? – M Metra

- How are the US guidelines developed? P Heidenreich
- How are the Canadian guidelines developed? J Ezekowitz
- How to reconcile new FDA and CMS pathways with guideline writing B Abraham
- Discussants: G Filippatos, N Stockbridge, B Zuckerman, M Fiuzat

15:10–15:55: Session 2 – Issues of Diagnosis

Chairs: S Anker / W Abraham / B Bozkurt

- The 3 "classes" of HF: HFrEF, HFmrEF & HFpEF all good? M Metra
- LVEF, does it still matter to differentiate groups of HFrEF patients? J Teerlink
- General Discussion
- Pro-Hormone biomarkers for prognosis & diagnosis of HF E Giannitsis
- What about ID assessment, now for all? E Jankowska
- General Discussion

15:55–16:35 Session 3 – AF issues

Chairs: J Butler / A. Coats

- The evidence for anti-coagulation J Cleland
- AF treatment in patients in HF What do different GLs say? P Kirchhof

----- 16:35-16:45: BREAK -----

- AF treatment in patients in HF The US perspective J Piccini
- CRT for patients with AF and HF G Hindricks
- General Discussion

16:45–18:00: Session 4 – SGLT2 inhibitors

Chairs: S Anker / J Butler

- For prevention: 5 drugs at 1A L Lund
- In diabetic HF patients 3 drugs at 1A G Filippatos
- In HFrEF regardless of diabetes 2 drugs at 1A R de Boer
- What can HF guidelines learn from CKD studies with SGLT2 inhibitors M Petrie
- General Discussion
- What about HFpEF: Is a GL-update already needed? A Coats
- What does HFA plan for this G Rosano
- What about SGLT2 inhibitors and changes in quality of life? Are separate recommendations needed? J Butler
- General Discussion

18:00–19:00: Session 5 - Device for HF therapy

Chairs: W Abraham / J Butler

- Neuromodulation F Zannad
- CCM: New Developments J Butler
- Intra-atrial shunt devices (covering all three approaches) W Abraham
- General Discussion
- General Discussion

19:00–19:30: Ad hoc Q & A session (part 1)

Program





OCTOBER 13th, 2021 - 13:30-18:00 CEST

13:30: Welcome A. Coats

13:35–14:20: Session 6 – Acute and severe HF related issues

Chairs: PC Schulze / O Chioncel

- Can (or should) GLs recommend drugs before regulatory approval, and if so, how?
 The vericiguat case B. Pieske
- Starting all HFrEF drugs in hospital? P Ponikowski
- Do Guidelines need to define up-titration steps and speed? I Piña
- What about Omecamtiv Mecarbil? J Teerlink
- · General Discussion

14:20–14:40: Session 7 – Co-morbidities and enablement strategies part 1 (obesity)

Chairs: J Butler / B Pieske

- Weight management in HF to improve exercise capacity with GLP1 agonists? M Packer
- Weight management in HFpEF to prevent hard outcomes J Butler General Discussion

14:40–15:15: Session 8 – Co-morbidities and enablement strategies part 2 (HK & ID)

Chairs: J Butler / S Zieroth

- Treatment of hyperkalemia M Kosiborod
- ID: The European Perspective P Ponikowski
- ID / Anemia: The US Perspective A Hernandez
- · General Discussion

15:15–15:50: Session 9 – Device for HF therapy part 1: focus on valvular heart disease

Chairs: W Abraham / S Anker

• Targeting FMR: MitraClip - W Abraham

- Targeting FMR: Carillon St von Bardeleben
- Targeting TR: (Edwards / Pascal system) J Hausleiter
- General Discussion

----- 15:50-16:00: BREAK -----

16:00–16:50: Session 10 – Controversial issues about ESC guidelines

Chairs: S Anker / PC Schulze / A Coats

GUIDED GROUP DISCUSSION

- Why 1B for entresto?
- Entresto after ACE inhibitors or straight away?
- 2B for Vericiguat, really?
- Why not class 1 for drugs that improve QoL & functional status more reliable than exercise?
- Is 1A for sotaglifozin in diabetic heart failure patients a surprise?
- Dual SGLT1 and SGLT2 inhibition: does it matter?
- The original "CardioMems restricted to LVEF ≤35%" statement and how it got correct in the printed version.

16:50–17:30: Session 11 – Open issues

Chairs: S Anker / P Ponikowski

GUIDED GROUP DISCUSSION

- Dual SGLT1 and SGLT2 inhibition: does it matter? D Bhatt
- Personalised approach and patients profiles when do you give ivabradine?
- HFmEF: now 2B for subgroups?!
- Finerenone and its success in DKD -> something for HF already?
- Separate QoL recommmendations for SGLT2 inhibitors? (the Preserved-HF case)
- Do we need a phenotyping approach for HFpEF as well?

17:30–18:00: Ad hoc Q & A session (part 2)

Conclusion: S Anker

---- In-person & Remote -----

Program





NOVEMBER 19th, 2021 - 15:30-19:00 CET

15:30–15:35: Welcome and summary of past discussions

15:35 -16:40: Session 12 – New HF trial results

Chairs: S Anker / J Butler

• GALACTIC-HF - J Teerlink

• AFFIRM-HF and more on Iron - P Ponikowski

• EMPEROR-Preserved – J Butler

• News on AF in HF – M Vaduganathan

• News on PRESERVED-HF - M Kosiborod

• News on EMPULSE - A Voors

• General Discussion

16:40 –16:55: Ad hoc Q & A session (part 3)

16:55–17:10: Session 13 – Revisiting possible treatment algorithms after the AHA (part 1)

GUIDED GROUP DISCUSSION

• Acute HF – all faculty

General Discussion

17:10–17:35: Session 14 – Revisiting possible treatment algorithms after the AHA (part 2)

Chairs: J Butler / A Coats

GUIDED GROUP DISCUSSION

• HFrEF – M&M oriented – all faculty

• HFrEF – symptoms – all faculty

• General Discussion

----- 17:35-17:45: BREAK -----

17:45–18:10: Session 15 – Revisiting possible treatment algorithms after the AHA (part 3)

Chairs: S Anker / J Butler

• HFpEF & HFmrEF - all faculty

General Discussion

18:10–18:40: Session 16 – Guidelines implementation and International cascade guidelines

Chairs: S Anker / A Coats

18:40–18:55: Ad hoc Q & A session (part 3)

18:55–19:00: Conclusions & End of Meeting



Information





Chairs (



Prof. Javed Butler

MD, MPH, MBA
Professor and Chairman of the
Department of Medicine at
the University of Mississippi in
Jackson, Mississippi, USA, Patrick
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Prof. William Abraham

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Prof. Stefan Anker

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Prof. Andrew Coats

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Faculty •

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R de Boer
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E Giannitsis
G Hindricks

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