

2025 POST-ESC HIGHLIGHT SYMPOSIUM (台北場)

時間: 114 年 9 月 14 日(星期日) 14:00-17:00

地點: 張榮發基金會國際會議中心 6 樓 602 會議室 (台北市中正區中山南路 11 號)

	Topic	Presenter	Moderator
14:00 – 14:05	Welcome Remarks		李貽恒
14:05 – 14:25	CAD and ACS	蘇峻弘	李貽恒
14:25 – 14:45	Arrhythmia and Device Therapy	李政鴻	林彥璋
14:45 – 15:05	Valvular Heart Disease & 2025 ESC guidelines	宋思賢	高憲立
15:05 – 15:25	Panel Discussion		陳文鍾
15:25 – 15:35	Healthy Break		
15:35 – 15:55	Pulmonary artery hypertension	吳俊賢	林彥宏
15:55 – 16:15	Lipid & 2025 ESC guidelines (Focused Update)	黃金洲	葉宏一
16:15 – 16:35	Hot Line and Late-Breaking Clinical Trials	林宗憲	謝宜璋
16:35 – 16:55	Panel Discussion		侯嘉殷
16:55 – 17:00	Closing Remarks		侯嘉殷

蘇峻弘

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不只好發老年族群！這些族群須小心「冠心病」危機

李政鴻醫師

主治專長

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重要著作：	Sung SH, Huang CJ, Cheng HM, Huang WM, Yu WC, Chen CH. Effect of Acoustic Cardiology-guided Management on 1-year Outcomes in

[Patients With Acute Heart Failure.](#) J Card Fail. 2020;(2):142-150.

Lee CW, Frerker C, Huang WM, Tsai YL, Huang CJ, Yu WC, Hsu CP, Chiang CE, Chen CH, **Sung SH.** [Feasibility and rationale of direct current cardioversion immediately after transcatheter percutaneous edge-to-edge mitral valve repair.](#) Eur J Clin Invest. 2020;50(10):e13274.

Chang HC, Huang WM, Yu WC, Cheng HM, Guo CY, Chiang CE, Chen CH, **Sung SH.** [Prognostic Role of Pulmonary Function in Patients With Heart Failure With Reduced Ejection Fraction.](#) J Am Heart Assoc. 2022;11(7):e023422.

Chang HC, Wei TW, Wu PY, Tsai MD, Yu WC, Chen CH, **Sung SH.** [TIFA protein expression is associated with pulmonary arterial hypertension.](#) Sci Rep. 2021;11(1):14140.

Chang HC, Huang CJ, Yang AC, Cheng HM, Chuang SY, Yu WC, Chiang CE, Chen CH, **Sung SH.** [Role of Heart Rate Variability in Association Between Glomerular Hyperfiltration and All-Cause Mortality.](#) J Am Heart Assoc. 2021;10(24):e021585.

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成人心臟學
心臟急重症加護
肺動脈高壓

專業執照

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黃金洲醫師

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	<p>2022: 第 18 屆榮總台灣聯合大學合作研究計畫『優良論文』第三名</p> <p>2022: 中華民國血脂及動脈硬化學會 111 年度海報論文獎第一名</p> <p>2024: 未來科技獎：「人工智慧輔助醫院血壓量測」</p>
重要著作：	<p>Hung MH, Shih LC, Wang YC, Leu HB, Huang PH, Wu TC, Lin SJ, Pan WH, Chen JW, <u>Huang CC</u>*.</p> <p>Prediction of masked hypertension and masked uncontrolled hypertension using machine learning. <i>Front Cardiovasc Med.</i> 2021;8:778306.</p> <p>Shih LC, Wang YC, Hung MH, Cheng H, Shiao YC, Tseng YH, <u>Huang CC</u>*, Lin SJ, Chen JW. Prediction of white-coat hypertension and white-coat uncontrolled hypertension using machine learning algorithm. <i>Eur Heart J Digit Health.</i> 2022;3:559-569.</p> <p><u>Huang CC</u>, Niu DM, Charng MJ. Genetic analysis in a Taiwanese cohort of 750 index patients with clinically diagnosed familial</p>

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YJ, **Huang CC***. The impacts of anemia burden on

clinical outcomes in patients with out-of-hospital

cardiac arrest. *Clin Cardiol.* 2024;47:e24175.

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CAD and ACS

蘇峻弘

For many years, patients with diabetes and no known coronary artery disease (CAD) were thought to have the same risk for future myocardial infarction (MI) as did patients with known CAD. CAD is a major determinant of the long-term prognosis among patients with T2DM. Furthermore, in patients with T2DM there is an increased mortality after MI, and worse overall prognosis in T2DM patients with CAD. In the past prior to 2008, the available drugs for glycemic control in DM had been largely neutral or even had some harmful effects. The newer agents, like SGLT-2 inhibitors and GLP-1 receptor agonists have recently been shown to be not only effective and safe for glycemic control, but also have cardio-protective effects. It is well recognized that as many as two-thirds of T2DM patients with either ACS or stable CAD have either previously diagnosed DM or will be subsequently diagnosed with it. It is, therefore, essential for the clinical cardiologist to be familiar with the latest therapeutic strategies and advances for the management of these patients. Given that there has been significant evolution in the development of pharmacologic management of T2DM patients and selection of the optimal antidiabetic strategy for T2DM patients with CAD is crucial. The TSOc consensus suggests the target of HbA1c <7%. Metformin remains the first-line therapy in diabetic patients with CAD, mainly based on the findings from the UKPDS trial, 3 meta-analyses,¹ observational study, and its effect on the reduction in CAC severity. For dual therapy, we recommend metformin plus SGLT-2 inhibitors, followed by metformin plus GLP-1 RAs, and then metformin plus TZDs (pioglitazone only). The PROactive trial, an important meta-analysis,⁷⁴ and 2 image studies (CHICAGO and PERISCOPE provided evidences to support the place of pioglitazone in the management of type 2 diabetes and CAD. The EMPA-REG OUTCOME trial, the CANVAS program, DECLARE TIMI-58 studies and the CVD-REAL Nordic study gave a rationale for the use of SGLT-2 inhibitors. Reassuringly, each of the completed large-scale 7 CVOTs and their recent meta-analysis of these trials demonstrates cardioprotective effect of GLP-1RA gave a rationale for the use of GLP-1 RAs. If the fourth drug is to be added, DPP-4 inhibitors are recommended due to their neutral effects and safety. Sulfonylurea did not have any positive trial to support its use, and the result of a Taiwanese cohort showed a worse outcome. In addition, the risk of hypoglycemia is well-known. Glinides and acarbose have low priority due to lack of any supporting evidence.

Arrhythmia and Device Therapy

李政鴻

Anderson-Fabry disease (AFD) is a rare X-linked inherited metabolic disorder which results in a deficiency or absence of the enzyme α -galactosidase A, leading to the accumulation of glycosphingolipids in various cells and organs, including the heart. Cardiac involvement is common, usually manifests as left ventricular hypertrophy (LVH), increased myocardial inflammation, myocardial fibrosis, heart failure and arrhythmias. Echocardiography and cardiovascular magnetic resonance (CMR) imaging offer distinctive and complementary use to assist in the diagnosis of AFD, including detection of the AFD cardiac phenotype, as well as the differentiation from other forms of LVH. Furthermore, CMR imaging - as the advanced cardiac imaging - holds promise in subclinical detection of AFD-related abnormalities as well as disease staging and prognostication.

In this presentation, we are going to demonstrate the features of AFD on echocardiography, as well as the characteristics on various imaging sequences for differential diagnosis in variable forms of LVH on CMR imaging.

Anemia is an important and common comorbidity in patients with heart failure. Patients with anemia and heart failure are associated with poor clinical status and worse outcomes than those without. Whether anemia is just a marker of heart failure severity or it is also involved in heart failure progression and outcomes and therefore should be treated is not entirely clear. Using erythropoiesis-stimulating agents to treat anemia in patients with heart failure has been evaluated intensively during the past several years and is regarded as a promising treatment strategy for heart failure patients. Unfortunately, it has been demonstrated in a large scale randomized controlled trial that these agents did not improve outcomes but were associated with a higher risk of adverse events. The iron deficiency in patients with heart failure can be absolute or functional. The former refers to that total body iron is decreased, and the latter is caused when total body iron is normal or increased but is inadequate to meet the needs of target tissues because of sequestration in the storage pool. It seems appropriate to supplement iron in patients with anemia resulting from absolute iron deficiency; however, it has been unclear whether and how absolute or functional iron deficiency should be treated in nonanemic patients with heart failure. Recently, some beneficial effects have been observed in small studies by administering intravenous iron in patients with heart failure and absolute or functional iron deficiency with or without anemia improves symptoms and exercise capacity. Nonetheless, their long-term outcomes and safety data are not yet available. In this lecture, we discuss the causes and pathogenesis of and treatment options for anemia and iron deficiency in patients with heart failure.

Pulmonary artery hypertension

吳俊賢

1. **Vericiguat**, a novel oral soluble guanylate cyclase stimulator, in this phase 3, randomized, double-blind, placebo-controlled trial showed beneficial effects in 5050 patients with NYHA II-IV and an EF < 45%. Vericiguat reduced 10% of composite endpoints including cardiovascular death and first heart failure hospitalization in mean 10.8 months.
2. **PARALLAX**, a prospective, randomized, controlled, double-blind multicentre clinical trial in patients with chronic symptomatic HF with EF >40%, New York Heart Association class II-IV symptoms, found sacubitril/valsartan reduced NT-proBNP levels at 12 weeks, but didn't increased 6-minutes walk distance at 24 weeks, compared to inhibitors of the RAS including ACE inhibitors or ARBs.
3. **EXPLORER-HCM**, a phase 3, randomised, double-blind, placebo-controlled trial found mavacamten (a first-in-class cardiac myosin inhibitor) improved exercise capacity, LVOT obstruction, NYHA functional class, and health status in patients with obstructive hypertrophic cardiomyopathy. Patients on mavacamten had greater reductions than those on placebo in post-exercise LVOT gradient (−36 mm Hg, 95% CI −43.2 to −28.1; $p<0.0001$). Thirty four% more patients in the mavacamten group improved by at least one NYHA class (80 of 123 patients in the mavacamten group vs 40 of 128 patients in the placebo group; 95% CI 22.2 to 45.4; $p<0.0001$).
4. **EMPEROR-Reduced**, a double-blind trial of 3730 patients with class II-IV HF and LVEF<40% to receive empagliflozin or placebo in addition to recommended therapy, found 25% reduction of cardiovascular death and HF hospitalization. The effect of empagliflozin on the primary outcome was consistent in patients regardless of the presence or absence of diabetes. The annual rate of decline in eGFR was slower in the empagliflozin group than in the placebo group, and empagliflozin-treated patients had a lower risk of serious renal outcomes.

Lipid & 2025 ESC guidelines

黃金洲

In the ESC 2025 conference, there was no ground-breaking news in the arrhythmic field. Nonetheless, there were a few interesting researches helping to fine tune the best strategy to treat patients in clinical practice.

As for the management of atrial fibrillation, the launch of ESC 2025 guideline did not differ much from previous one. The low risk of stroke in women with no other risk factor was incorporated into the guideline and needed other risk factors to warrant anticoagulation. The coverage of device detected atrial high rate was new and recommendation of anticoagulation was put into the guideline. However, more evidence is needed. The use of aspirin for stroke prevention in AF patients is totally out of the recommendation due to the similar bleeding risk and less efficacy of stroke prevention than warfarin.

To clarify the role of ablation in treating atrial fibrillation, there are many registries going on all over the world. The one year recurrent rate was around 70% among all the registries, even with the better ablation tools and the improvement of the durability of PV isolation, implying that 30% of the AF patients had triggering foci outside PVs.

There were some post-marketing registries of NOAC, aiming to study the safety of NOAC in real world, including Norway registry, ORBIT-AF registry, and XaPASS. The results were generally consistent with the phase III RCTs. Edoxaban was mostly not in the analysis of these registries due to its late launch. Its study, ENSURE-AF, showed the safety profile of edoxaban in the use of electrical cardioversion. As for the anti-dote of Xa inhibitor, ANNEXA-4 revealed the efficacy and safety of the drug, andexanet, in reversing the effect of apixaban or revaloxaban.

As for ventricular tachyarrhythmia, DANISH proved no survival benefit of ICD in patients with non-ischemic cardiomyopathy as primary prevention. Numerous studies searched for better risk stratification markers, but the adoption for clinical use needs further studies. The same scenario applied to ischemic cardiomyopathy too. The hot topic of HRS this year, early repolarization, was not hot anymore in ESC. However, it seems to be a good marker of future ventricular arrhythmia events in heart failure patients.

Hot Line and Late-Breaking Clinical Trials

林宗憲

Many clinical trials are presented in the “Hot-Line sessions” of European Society of Cardiology scientific conference this year. Studies, including DANISH and REM-HF for the efficacy of implantable cardioverter defibrillator in heart failure patients were presented and discussed. Strategies of dual antiplatelet treatment (DAPT) and platelet function monitoring (ANTARCTIC) challenged the current concept of DAPT treatment. Management for hyperlipidemia is still a hot topic in this year. Study from Japan (IJ-PROPER) argued the effect of ezetimibe on acute coronary event. ESCAPE trial investigated the effect of alirocumab, a PCSK9 inhibitor, on the frequency of lipoprotein apheresis in a randomized phase 3 setting. CE-MARC 2 trial investigated the diagnostic accuracy for coronary artery disease by various image modality, including MR and SPECT and presented in the meeting. The choice of antiplatelet agents and its effects in patients without stenting were discussed in EROSION and PRAGUE-18 trials. ENSURE-AF study reported the effects of edoxaban vs. enoxaparin/warfarin in subjects undergoing cardioversion of atrial fibrillation. These results of clinical trials update the concept and provide information on treatment strategies on patients with cardiovascular diseases.