Prediabetes and insulin resistance in a population of patients with heart failure and reduced or preserved ejection fraction but without diabetes, overweight or hypertension

Tran Kim Son1, Ngo Hoang Toan1, Nguyen Thang2, Huynh Le Trong Tuong3, Hoang Anh Tien4, Nguyen Hai Thuy4, Huynh Van Minh4 and Paul Valensi5*

Abstract

Background: The relationships between glucose abnormalities, insulin resistance (IR) and heart failure (HF) are unclear, especially regarding to the HF type, i.e., HF with reduced (HFrEF) or preserved (HFpEF) ejection fraction. Overweight, diabetes and hypertension are potential contributors to IR in persons with HF. This study aimed to evaluate the prevalence of prediabetes and IR in a population of Vietnamese patients with HFrEF or HFpEF but no overweight, diabetes or hypertension, in comparison with healthy controls, and the relation between prediabetes or IR and HF severity.

Methods: We conducted a prospective cross-sectional observational study in 190 non-overweight normotensive HF patients (114 with HFrEF and 76 with HFpEF, 92.6% were ischemic HF, mean age was 70.1 years, mean BMI 19.7 kg/m²) without diabetes (neither known diabetes nor newly diagnosed by OGTT) and 95 healthy individuals (controls). Prediabetes was defined using 2006 WHO criteria. Glucose and insulin levels were measured fasting and 2 h after glucose challenge. IR was assessed using HOMA-IR and several other indexes.

Results: Compared to controls, HF patients had a higher prevalence of prediabetes (63.2% vs 22.1%) and IR (according to HOMA-IR, 55.3% vs 26.3%), higher HOMA-IR, insulin/glucose ratio after glucose and FIRI, and lower ISIT0 and ISIT120 (<0.0001 for all comparisons), with no difference for body weight, waist circumference, blood pressure and lipid parameters. Prediabetes was more prevalent (69.3% vs 53.9%, p=0.03) and HOMA-IR was higher (p<0.0001) in patients with HFrEF than with HFpEF. Among both HFrEF and HFpEF patients, those with prediabetes or IR had a more severe HF (higher NYHA functional class and NT-proBNP levels, lower ejection fraction; p=0.04–<0.0001) than their normoglycemic or non-insulinresistant counterparts, with no difference for blood pressure and lipid parameters.

Conclusion: In non-diabetic non-overweight normotensive patients with HF, the prevalence of prediabetes is higher with some trend to more severe IR in those with HFrEF than in those with HFpEF. Both prediabetes and IR are...
Background
Heart failure (HF) is a leading cause of morbidity and mortality in Western countries. Its prevalence has been increasing, as well as its financial burden [1–3]. In recent cardiovascular outcome trials including patients with type 2 diabetes and a history of cardiovascular disease or several associated risk factors, the incidence of hospitalization for HF was comparable to the incidence of acute myocardial infarction [4]. The prevalence of patients with both HF and diabetes keeps increasing as the population is ageing [5] and diabetes is among the most common co-morbidities in HF patients, potentially increasing the risk of hospitalization and death [5]. Interestingly, HF with preserved ejection fraction (HFpEF) is currently the most frequent form of HF [6].

Glucose abnormalities often remain undiagnosed in HF patients [7] although the incidence of diabetes is high [8]. An exploratory analysis from the Dapagliflozin and Prevention of Adverse Outcomes in Heart Failure (DAPA-HF) trial looked at the incidence of diabetes (HbA1c $\geq$ 6.5% or initiation of a glucose-lowering agent) in the subgroup of patients with HF and reduced left ventricle ejection fraction (HFrEF), no prior history of diabetes and an HbA1c level $< 6.5\%$ at baseline. In this subgroup, the incidence of newly-diagnosed diabetes was 7.1% in the placebo group over a median follow-up of 18 months [9].

Insulin resistance (IR) is frequent in HF patients, independently of an ischemic etiology [10, 11]. In a community cohort, IR was associated with incident HF and this relationship was not modified by BMI [12]. IR was also shown to be a predictor of death among HF patients without known diabetes [13]. IR has been frequently described among patients with HFrEF [13, 14]. Whether or not the prevalence of IR is different in patients with HFpEF has not been clearly established. A recent study on a small series showed that older patients (70–90 years) with either systolic HF or diastolic HF were more insulin resistant than same age healthy volunteers [14]. Another study suggested more severe IR in HFrEF patients than in HFpEF patients [15]. However, most studies looking at the association between HF and IR included few patients, and glucose status was not reported or patients with diabetes were not clearly excluded. While IR and HF may participate into a vicious circle, the precise relationship between IR and HF is an interesting but unsolved issue. In addition, obesity and hypertension are well-known risk factors for IR, type 2 diabetes but also HF [16, 17].

Thus, the relationships between glucose abnormalities, IR and HF need to be clarified. Known diabetes or newly diagnosed diabetes, obesity and hypertension should be considered as potential contributors to IR in the HF population. In addition, these relationships should be analyzed considering separately HFrEF and HFpEF.

The aim of this study was to evaluate the prevalence of prediabetes and IR in a population of non-diabetic non-overweight normotensive Vietnamese patients with HFrEF or HFpEF compared to healthy controls, and the relation of prediabetes and IR with HF severity.

Methods
Study design
We conducted a prospective cross-sectional observational study at Can Tho Central General Hospital, Vietnam, from April 2013 to May 2016, comparing consecutive patients with HF to healthy subjects.

This study was approved by the Institutional Review Board of this hospital. Information and data were secured. Patients and controls gave their consent for participation to the study. All the costs for lab tests were supported by the hospital research funding.

HF patients had been hospitalized for congestive episodes at least 6 months ago. Participants were in stable condition at the investigation time, with no change in treatments within this 6-months interval. The investigations were performed as outpatients or during hospitalization for clinical assessment and drug delivery. IR assessment and other biochemical measurements were performed a few days before echocardiography.

Healthy subjects were volunteers whose tests were performed in the hospital during an annual health check-up.

Study population
Inclusion and exclusion criteria
Patients were included when they had a diagnosis of HF according to the 2016 European Society of Cardiology (ESC) guidelines [18]. The diagnosis of chronic HF was based on clinical symptoms and signs and confirmed with echocardiography showing evidence for an impaired
myocardial function. According to guidelines [18], HFrEF was defined as LVEF ≤ 40% and HFpEF as LVEF ≥ 50%.

Patients with LVEF between 40 and 50% were excluded. We also excluded all patients with disorders or treatments which might potentially alter insulin sensitivity: (i) known diabetes, hypertension, chronic kidney disease defined as eGFR < 90 ml/min/1.73 m², newly-diagnosed diabetes during Oral Glucose Tolerance Test (OGTT); (ii) BMI > 23 kg/m² (the threshold for overweight diagnosis in Asian populations) or waist circumference > 90 cm for men, > 80 cm for women [19]; (iii) current hormonal therapies (substitutive treatment for menopause, corticosteroids and beta-adrenoreceptor agonists).

Healthy controls were randomly selected by screening among healthy volunteers. They had similar age, gender, BMI, and other anthropometric characteristics and no history, symptoms or signs of cardiovascular disease including HF, no acute or chronic disease and no treatment susceptible to alter IR.

Clinical assessment
All HF participants underwent a clinical comprehensive evaluation including an interview, complete physical examination, and echocardiography. HF was considered of ischemic cause when the patient had been previously diagnosed with coronary artery disease with no other explanation for HF.

In HF and control subjects, waist circumference was measured at the end of mild expiration, parallel to the floor using a supported tape. Measurement was made naked trunk, at the narrowest section, half-distance between the lower border of the costal margin and the iliac crest. Body weight and height were obtained with light clothing on. Body mass index was calculated using the weight (kg)/height (m)² formula. Blood pressure was measured on both arms in the sitting position, and the diastolic pressure was measured by auscultation. Weight, height, and waist circumference were measured without underwear, and processed for biochemical measurements.

Biochemical measurements
Blood samples were collected after an overnight fast. They were immediately sent to the central hospital laboratory and processed for biochemical measurements.

Assessment of glucose status and insulin resistance
An OGTT was performed and plasma glucose and insulin levels were measured at fasting (G₀ and I₀) and 2 h after glucose intake (G₂ and I₂).

Three prediabetic conditions were considered, as defined by WHO 2006 criteria [20]: impaired fasting glucose (IFG) by G₀ = 6.1–6.9 mmol/l (110–125 mg/dl), impaired glucose tolerance (IGT) defined by G₂ = 7.8–11.1 mmol/l (140–200 mg/dl), or combined IFG and IGT. Normal glucose tolerance (NGT) was defined by G₀ < 6.1 mmol/l (110 mg/dl) and G₂ < 7.8 mmol/l (140 mg/dl).

Insulin resistance was assessed using several indexes: HOMA-IR index (HOMeostatic Model Assessment of Insulin Resistance: I₀ × G₀/22.55) [21] with a cut-off value defined by the highest quartile of the control group; QUICKI [QUantitative Insulin-sensitivity Check Index: 1/log(I₀ + G₀)] [22] with a cut-off value defined as the lowest quartile of the control group; and by I₀, I₀/ G₀, I₂ and I₂/G₂ with cut-off values defined the respective mean ± 1SD values of the control group. We also calculated additional insulin resistance indexes: fasting Insulin Sensitivity Index [ISIT0 = 10,000/(I₀ × G₀)], 2-h-Insulin Sensitivity Index [ISIT120 = 10,000/(I₂ × G₂)] [23] and Fasting Insulin Resistance Index [FIRI = (I₀ × G₀/25)] [24].

Analytical methods
Plasma glucose, HbA1c, total cholesterol, triglycerides, HDL-cholesterol, and insulin were measured using HumaStar 600 (Wiesbaden, Germany). LDL-cholesterol was calculated using Friedwald formula. NT-proBNP was measured by electrochemiluminescence immuno-assay using Cobas E (Roche Diagnostics, USA).

Statistical analyses
The normality of the data was assessed using the Kolmogorov–Smirnov test. Data are presented as mean ± standard deviation (SD) or as median [interquartile] if the distribution was not normal. Between-groups comparisons for continuous variables were performed by analyses of variance if normally distributed or by nonparametric tests if not. Between-groups comparisons for categorical parameters were performed by chi square tests. Odds ratios (OR) with 95% confidence intervals (CI) for the risk of IR in HF patients are reported. A p value < 0.05 was considered as statistically significant. Statistical analyses were carried out using SPSS statistical software version 22.0.

Results
Clinical characteristics of the study population
Among 245 HF patients fulfilling the selection criteria, 215 gave their consent for participation to the study. Among them, 25 were either diagnosed with diabetes during OGTT or withdrew consent. Thus, the study population included 190 HF patients. Age range was 50 to 85 years, 114 patients had HFrEF and 76 HFpEF (Fig. 1). Baseline characteristics of these patients and of the 95 control subjects are shown in Table 1. Sex-ratio, age, BMI and waist circumference did not differ significantly among healthy volunteers. They had similar age, gender, BMI, and other anthropometric characteristics and no history, symptoms or signs of cardiovascular disease including HF, no acute or chronic disease and no treatment susceptible to alter IR.
those with HFrEF and those with HFpEF. Total and LDL cholesterol were higher in HFrEF patients. Cardiac ischemic disease was the main cause of HF (92.6%).

Compared to HFpEF patients, HFrEF patients had higher NT-pro-BNP levels, were more often in NYHA functional class III or IV, and had more often HF from ischemic cause (Table 1).

Glycemic status
Based on OGTT results, the prevalence of prediabetes was significantly higher in HF patients compared to control subjects (63.2% vs 24.2%, respectively; p<0.0001). Among HF patients, the prevalence of IGT was significantly higher and the prevalence of NGT significantly lower (p<0.0001) in patients with HFrEF compared to those with HFpEF (Table 1).

Compared to control subjects, G0 and G2, and I0 and I2 were higher in HF patients. HbA1c levels did not differ significantly, and none of the HF patients had an HbA1c level ≥ 6.5%. G0 and G2 were higher in HFrEF compared to HFpEF patients, without significant difference for HbA1c levels (Table 1).

Prevalence of insulin resistance in patients with HFrEF and HFpEF
Most of the IR indexes differed significantly between HF patients and controls. HOMA-IR, I2/G2 and FIRI were higher, and ISIT0 and ISIT120 lower in HF patients compared to controls. Compared to HFpEF patients, HFrEF patients had higher HOMA-IR with a trend for higher FIRI and lower ISIT0 (Table 2).

We then considered the cut-off values of IR indexes on the basis of values in the control group. The cut-off values were 2.53 (highest quartile) for HOMA-IR index; 0.33 (lowest quartile) for QUICKI and the mean ±1SD for I0/G0 and I2/G2 indexes in the control group. The prevalence of IR as defined by values over the cut-off values according to HOMA-IR was significantly higher in patients with HFrEF and HFpEF (58.8% and 50.0%, respectively) than in the control group (p<0.001). Based on the other IR indexes including I0/G0, I2/G2 and QUICKI, the prevalence of IR in HFrEF and HFpEF groups was 30.7% to 60.5% and 26.3% to 50%, respectively, without significant difference between the two groups (Table 3).

Likelihood of IR in HF patients
Compared to controls, HFrEF patients had a highly significant likelihood of IR according to these cut-off values for all the indexes with very high OR, especially for HOMA index and QUICKI. The same significant trend (except for I2/G2) was observed in HFpEF patients. Meanwhile, there was no significant difference between HFrEF and HFpEF populations for the risk of IR based on any of these indexes (Table 4).
Table 1 Characteristics of the study population

|                           | HF patients | Controls | p* | p**  |
|---------------------------|-------------|----------|----|------|
|                           | Total HF    | HFrEF    | HFP EF | (n = 190) | (n = 114) | (n = 76) |       | (n = 95) |      |
| Gender (%)                |             |          |      |       |          |          |       |         |       |
| Male                      | 95 (50)     | 55 (48.2)| 40 (52.6) | 0.813 | 46 (48.4) | 0.802 |
| Female                    | 95 (50)     | 59 (51.8)| 36 (47.4) |       | 49 (51.6) |       |
| Age (years)               | 70.1 ± 14.5 | 70.5 ± 14.0 | 69.5 ± 15.4 | 0.162 | 66.7 ± 14.7 | 0.064 |
| BML (kg/m²)               | 19.7 ± 1.5  | 19.8 ± 1.4 | 19.5 ± 1.6  | 0.202 | 19.9 ± 2.0 | 0.091 |
| Waist circumference (cm)  | 74.4 ± 4.3  | 74.2 ± 4.2 | 74.6 ± 4.6  | 0.411 | 74.6 ± 4.3 | 0.743 |
| Systolic BP (mmHg)        | 116.5 ± 15.4 | 117.5 ± 15.8 | 115.0 ± 14.7 | 0.233 | 114.3 ± 12.0 | 0.223 |
| Diastolic BP (mmHg)       | 71.1 ± 10.4 | 71.3 ± 10.3 | 70.9 ± 10.7 | 0.646 | 71.7 ± 9.4 | 0.646 |
| LVEF (%)                  | 42.9 ± 14.4 | 32.2 ± 5.8 | 58.9 ± 6.3 | < 0.001 |          |       |
| HF ischemic etiology (%)  | 176 (92.6)  | 110 (96.5%) | 66 (86.8%) | 0.013 |          |       |
| NYHA class                |             |          |      |       |          |          |       |         |       |
| II                        | 32 (16.8)   | 9 (7.9)  | 23 (30.3) | < 0.0001 |          |       |
| III                       | 150 (78.9)  | 97 (85.1) | 53 (69.7) |       |          |       |
| IV                        | 8 (4.2)     | 8 (7.0)  | 0 (0)    |       |          |       |
| NT-pro-BNP (pg/ml)        | 9931 ± 7747 | 11,550 ± 8702 | 7504 ± 5215 | < 0.0001 |          |       |
| HbA1c (%)                 | 5.70 ± 0.50 | 5.71 ± 0.50 | 5.70 ± 0.51 | 0.81 |          |       |
| HbA1c ≥ 6.5% (%)          | 0           | 0        | 0        |       |          |       |
| G0 (mmol/l)               | 5.28 ± 0.79 | 5.37 ± 0.76 | 5.13 ± 0.82 | < 0.0001 | 4.81 ± 0.81 | < 0.0001 |
| G2 (mmol/l)               | 8.07 ± 1.18 | 7.83 ± 1.14 | 6.93 ± 0.93 | < 0.0001 | 7.13 ± 1.73 | < 0.0001 |
| I0 (µU/ml)                | 12.1 [5.3–23.7] | 12.6 [5.5–26.4] | 11.1 [4.5–21.7] | 0.675 | 8.2 [5.1–13.0] | 0.003 |
| I2 (µU/ml)                | 90.3 [47.6–136.6] | 95.0 [45.3–141.2] | 73.6 [48.0–127.0] | 0.054 | 50.0 [24.0–96.0] | < 0.0001 |
| Total cholesterol (mmol/l)| 4.17 ± 1.50 | 4.44 ± 1.66 | 3.77 ± 1.13 | 0.03 |          |       |
| Triglycerides (mmol/l)    | 1.57 ± 1.13 | 1.74 ± 1.33 | 1.32 ± 0.64 | 0.13 |          |       |
| LDL-cholesterol (mmol/l)  | 2.62 ± 0.98 | 2.78 ± 1.04 | 2.38 ± 0.82 | 0.05 |          |       |
| HDL-cholesterol (mmol/l)  | 1.01 ± 0.30 | 1.04 ± 0.31 | 0.97 ± 0.28 | 0.113 |          |       |
| Glycemic status (%)       |             |          |      |       |          |         | <0.0001 |       |
| IFG alone                 | 4 (2.1)     | 4 (2.1)  | 0       | 0.032 | 6 (6.3)  | < 0.0001 |
| IGT alone                 | 94 (49.5)   | 64 (56.1) | 30 (39.5) | 15 (15.8) |          |       |
| IGT + IFG                 | 22 (11.6)   | 11 (9.6)  | 11 (14.5) | 2 (2.1) |          |       |
| NGT                       | 70 (36.8)   | 35 (30.7) | 35 (46.1) | 72 (75.8) |          |       |
| Medications (%)           |             |          |      |       |          |         | <0.0001 |       |
| ACEIs/ARBs                | 88 (46.3)   | 70 (61.4) | 18 (23.7) | < 0.0001 |          |       |
| Aldosterone antagonists   | 61 (32.1)   | 61 (53.5) | 0 (0)   | < 0.0001 |          |       |
| Beta-blockers             | 120 (63.2)  | 91 (79.8) | 29 (38.2) | < 0.0001 |          |       |
| Diuretics                 | 52 (27.4)   | 46 (40.4) | 6 (7.9)  | < 0.0001 |          |       |
| Ivabradine                | 40 (21.1)   | 36 (31.6) | 4 (5.3)  | < 0.0001 |          |       |
| Digoxin                   | 13 (6.8)    | 13 (11.4) | 0 (0)   | 0.002 |          |       |
| Antiplatelets             | 176 (92.6)  | 110 (96.5) | 66 (86.8) | 0.014 |          |       |
| Statin                    | 159 (83.7)  | 98 (86.0) | 61 (80.3) | 0.310 |          |       |
| Nitrate                   | 31 (16.3)   | 19 (16.7) | 12 (15.8) | 0.950 |          |       |

ACEIs/ARBs: angiotensin-converting-enzyme inhibitors/angiotensin II receptor blockers; BP: blood pressure; HF: heart failure; HFrEF and HFP EF: HF with reduced and preserved ejection fraction; BMI: body mass index; G0 and I0: plasma glucose and insulin levels measured fasting; G2 and I2: plasma glucose and insulin levels measured 2 h after glucose intake; IFG: impaired fasting glucose; IGT: impaired glucose tolerance; NGT: normal glucose tolerance; NT-pro-BNP: N-terminal pro Brain Natriuretic Peptide; NYHA: New-York Heart Association

*p values for comparisons between HFrEF and HFP EF patients
**p for comparisons between the total HF group and the control group
Comparison between HF patients with or without prediabetes and with or without IR

Among HFrEF and HFpEF patients considered separately, those with prediabetes were more often in high NYHA class and had higher NT-proBNP level and lower LVEF than those with NGT. IR was also more pronounced in those patients as shown by significantly different values for all the IR indexes and higher glycemic levels, with no other difference for clinical or biological parameters nor for the etiology of HF (Table 5). Similar trends were found in patients with IR (as defined by HOMA-IR ≥ 2.53) compared to those without, with no significant difference for BMI, waist circumference and lipid parameters. Nevertheless, more
patients with dysglycemia were found among patients with IR (Table 6).

**Discussion**

The most important findings of this study are that in a Vietnamese population of patients with HF and no diabetes, overweight or hypertension, (i) the prevalence of prediabetes and IR is high, (ii) prediabetes and IR are associated with a higher degree of HF, (iii) both findings affect similarly HFrEF and HFpEF patients.

Prediabetes, an underdiagnosed condition and a marker of worse cardiac function in patients with HFrEF or HFpEF It is commonly agreed that 20–30% of patients with HFrEF have had a previous diagnosis of diabetes. European guidelines on diabetes, prediabetes and
Table 6  Comparisons of HFrEF and HFpEF patients by insulin resistance status, i.e., with HOMA ≥ 2.53 or < 2.53

| Gender (%)          | HOMA ≥ 2.53 | HOMA < 2.53 | p values | HOMA ≥ 2.53 | HOMA < 2.53 | p values |
|---------------------|-------------|-------------|----------|-------------|-------------|----------|
| Gender (%)          | HOMA ≥ 2.53 | HOMA < 2.53 |          | HOMA ≥ 2.53 | HOMA < 2.53 |          |
| Male                | (n = 67)    | (n = 47)    | 0.614    | (n = 38)    | (n = 38)    | 0.168    |
| Female              |             |             |          |             |             |         |
| Age (years)         | 70.3 ± 13.2 | 70.8 ± 15.1 | 0.849    | 71.1 ± 13.7 | 67.9 ± 17.0 | 0.367    |
| BMI (kg/m²)         | 19.8 ± 1.5  | 19.8 ± 1.1  | 0.915    | 19.7 ± 1.6  | 19.3 ± 1.7  | 0.294    |
| Waist circumference (cm) | 74.7 ± 3.9 | 73.5 ± 4.5 | 0.138 | 75.0 ± 4.5 | 74.2 ± 4.7 | 0.441 |
| Systolic BP (mmHg)  | 116.9 ± 14.4 | 118.3 ± 17.7 | 0.661 | 118.3 ± 13.1 | 111.8 ± 15.6 | 0.052 |
| Diastolic BP (mmHg) | 71.1 ± 9.2  | 71.5 ± 11.8 | 0.817    | 73.5 ± 9.59 | 68.3 ± 11.3 | 0.034    |
| LVEF (%)            | 29.6 ± 5.9  | 36.0 ± 3.1  | < 0.0001 | 54.5 ± 3.2  | 63.3 ± 5.5  | < 0.0001 |
| NYHA class          |             |             |          |             |             |         |
| II                  | 1 (1.5)     | 8 (17.0)    | < 0.001  | 3 (7.9)     | 20 (52.6)   | < 0.0001 |
| III                 | 58 (86.6)   | 39 (83.0)   |          | 35 (92.1)   | 18 (47.4)   |          |
| IV                  | 8 (11.9)    | 0 (0)       |          | 0 (0)       | 0           |          |
| NT-pro-BNP (pg/ml)  | 15,046 ± 9104 | 6656 ± 4891 | < 0.0001 | 10,857 ± 4823 | 4151 ± 2962 | < 0.0001 |
| HbA1c (%)           | 5.68 ± 0.51 | 5.75 ± 0.49 | 0.487    | 5.66 ± 0.56 | 5.72 ± 0.48 | 0.588    |
| Total cholesterol (mmol/l) | 4.44 ± 1.25 | 4.44 ± 2.14 | 0.995    | 3.80 ± 1.07 | 3.76 ± 1.20 | 0.88     |
| Triglycerides (mmol/l) | 1.89 ± 1.33 | 1.53 ± 1.34 | 0.163    | 1.46 ± 0.71 | 1.19 ± 0.55 | 0.075    |
| LDL-cholesterol (mmol/l) | 2.78 ± 0.89 | 2.80 ± 1.25 | 0.937    | 2.44 ± 0.77 | 2.31 ± 0.89 | 0.518    |
| HDL-cholesterol (mmol/l) | 1.01 ± 0.30 | 1.10 ± 0.32 | 0.214    | 0.97 ± 0.29 | 0.98 ± 0.29 | 0.937    |

Glycemic status

| NGT, n (%)          | HOMA ≥ 2.53 | HOMA < 2.53 | p values | HOMA ≥ 2.53 | HOMA < 2.53 | p values |
|---------------------|-------------|-------------|----------|-------------|-------------|----------|
| II                  | 7 (10.4)    | 28 (59.6)   | < 0.0001 | 12 (31.6)   | 23 (60.5)   | 0.111    |
| IGT, n (%)          | 49 (73.1)   | 15 (31.9)   |          | 19 (50)     | 11 (28.9)   |          |
| IFG, n (%)          | 2 (3.0)     | 4 (4.3)     | 0        | 0           | 0           |          |
| IGT + IFG, n (%)    | 9 (13.4)    | 2 (4.3)     | 0.04     | 7 (18.4)    | 4 (10.5)    |          |

BMI body mass index; BP blood pressure; FRII fasting insulin resistance index; G0 and G1 plasma glucose and insulin levels measured fasting; G2 and G3 plasma glucose and insulin levels measured 2 h after glucose intake; HFrEF and HFpEF HF with reduced and preserved ejection fraction; HOMA-IR homeostatic model assessment of insulin resistance; IFG impaired fasting glucose; IGT impaired glucose tolerance; ISIT0 fasting insulin sensitivity index; ISIT120 2 h-insulin sensitivity index; NGT normal glucose tolerance; NT-pro-BNP N-terminal pro Brain Natriuretic Peptide; NYHA New York Heart Association; QUICKI quantitative insulin-sensitivity check index

cardiovascular disease recommend performing an OGTT in all patients with CVD and no known diabetes [25]. This recommendation is supported by data in patients with coronary disease [26]. In addition, the presence of prediabetes is associated with an increased risk of cardiovascular events and incident type 2 diabetes. In a series of stable subjects with prediabetes who underwent elective coronary angiography, the presence of subclinical myocardial necrosis as detected by high-sensitivity cardiac troponin T was reported to be prevalent and portend heightened long-term adverse cardiovascular event risk [27]. Thus, this marker may help to stratify cardiovascular risk in prediabetic subjects. Regarding the risk of HF, in the Multi-Ethnic Study of Atherosclerosis (MESA) HbA1c and fasting plasma glucose in the diabetes but not in the prediabetes range were each associated with...
higher risks of incident hospitalization for HF (HFrEF or HFrEF) [28]. Stratifying the risk of incident type 2 diabetes in prediabetic subjects using new markers [29] may also be helpful.

The reports on the frequency of glucose abnormalities in non-ischemic HF patients are scarce and included only a small number of participants [30, 31]. In a series of 227 HFrEF patients who had an OGTT, 23% were classified as having IGT and 18% as having newly diagnosed diabetes, and similar percentages were found in patients with either ischemic or non-ischemic HF [7]. In a subanalysis of the SUPPORT trial including 535 HF patients without known diabetes (18% with HFrEF and 62% with HFrEF), IGT was found in 23% of those participants [32]. However, there was no published study comparing adequately the prevalence of newly diagnosed glucose disorders in HFrEF and HFrEF patients. In our study, we were careful to exclude not only patients with previously diagnosed diabetes but also those diagnosed with diabetes according to OGTT and HbA1c was <6.5% in all the participants. Furthermore, we excluded potential confounders such as overweight, hypertension, impaired renal function and current hormonal therapies that could have changed glucose metabolism. Compared with controls who had the same sex-ratio and BMI but were slightly younger, the prevalence of prediabetes (mostly IGT) during OGTT was more than doubled, reaching 63.2%, in HF participants. Noteworthy, if only fasting plasma glucose levels had been measured, 94 participants with IGT alone out of the 120 patients with prediabetes (78%) would have been undiagnosed.

Prediabetes affected slightly more patients with HFrEF than those with HFrEF (66% vs 54% respectively). When HFrEF and HFrEF patients were taken separately, those with prediabetes had more severe HF with a higher NYHA functional class and higher plasma NT-proBNP levels. Ultrasound-measured LVEF was lower in participants with prediabetes compared to their normoglycemic counterparts while age, gender, BMI, waist circumference, blood pressure, lipid levels and the proportion of ischemic HF did not differ significantly.

In HF patients, a subgroup analysis of CHARM study reported a remarkably high prevalence of dysglycemia as detected by HbA1c ≥6.0% irrespective of ejection fraction phenotype and an association between dysglycemia and a higher risk of adverse clinical outcomes [33]. However, HbA1c ≥6.0% is not currently recognized as a cut-off point for dysglycemia and does not allow a clear diagnosis of glycemic status. The survival curve for HF patients with IGT or newly-diagnosed diabetes on OGTT was reported to be in intermediate positions between the curve for NGT and the one for previously diagnosed diabetes, with similar mortality rates for ischemic and non-ischemic HF [7]. In line with that study, our results showing that prediabetes is associated with worse cardiac function in HFrEF and HFrEF highlights the use of OGTT in risk stratification. Further studies should investigate whether or not interventions in patients with prediabetes and more specifically with IGT, may slow or reverse HF progression and related outcomes.

**Insulin resistance, a highly prevalent disorder associated with worse cardiac function in patients with HFrEF or HFrEF**

In our study, various indexes were used to assess IR and IR was more pronounced in HF patients compared to healthy controls with a trend for more pronounced IR in HFrEF compared to HFrEF patients. When considering the cut-off values of IR indexes in the control group, the prevalence of IR was doubled in HF patients compared to controls. However, the prevalence of IR was close in both HF groups (58.8% and 50% according to HOMA-IR index, respectively) with no significant difference between HF groups for the risk of IR using several IR indexes. Our results are in line with previous studies reporting a high prevalence of IR among often overweight patients with HFrEF [13, 14, 34]. Noteworthy, cardiac ischemic disease was form far the most frequent cause of HF (92.6%) in our population, and that could explain, at least partly, the high prevalence of prediabetes and IR we report. Only a few publications compared IR in HFrEF and HFrEF patients and usually on a limited number of patients. Our results are consistent with two previous studies showing no statistically significant difference for HOMA-IR in HFrEF and HFrEF patients [14, 34]. In a recent study including only 60 HF patients, fasting and dynamic measures of IR were performed in non-diabetic individuals with stable ambulatory HFrEF or HFrEF and showed more severe IR in patients with HFrEF on the dynamic tests [15]. Importantly, in our study, the high prevalence of IR was shown in both HFrEF and HFrEF patients and for the first time after exclusion not only of the patients with previously diagnosed diabetes but also of those with newly diagnosed diabetes on OGTT and after exclusion of patients with overweight or hypertension and of other confounding factors. Furthermore, while most reports used only the HOMA-IR index to assess IR, our study demonstrates that combining this index with other indirect indexes contributes to identify IR risk in patients with HF. Specifically, QUICKI index and I0/G0 were relevant parameters to identify this risk and hence their use could be suggested in IR assessment. Further studies in larger populations will be useful to precise incremental insights of IR differences between HFrEF and HFrEF patients.
We did not observe any significant difference in metabolic syndrome components according to the presence or absence of IR, except for dysglycemia which was more prevalent in participants with IR. Whether IR is a consequence, or a cause of HF is unclear. As IR was highly prevalent despite normal body weight and no excess in abdominal adiposity and was not associated with pronounced metabolic disorders, IR was more likely the consequence rather than the cause of HF and this might secondarily lead to dysglycemia. The high incidence of new diabetes in HF patients is consistent with this hypothesis [9]. However, the glycemic alterations induced by IR, ranging from mild dysglycemia to diabetes, may conversely increase the risk of HF and HF progression [5, 7]. How IR develops and worsens in HF patients is not well-understood, but IR may derive from many factors: enhanced sympathetic activation, loss of skeletal muscle mass, endothelial dysfunction, forced sedentary lifestyle due to reduced cardiac output and increased fatigability, a potential effect of increased circulating cytokines, and molecular mechanisms [6, 35–42]. The main mechanism involves probably 'neurohormone hypothesis' [43, 44]. Humoral neural stimuli are enhanced in both HFrEF and HFP EF, leading to an activation of the sympathetic nervous system and subsequently raised catecholamine levels. Catecholamine increase contributes to numerous adverse metabolic effects: negative impacts on insulin signaling and glucose utilization in skeletal muscle, reduction of insulin secretion, stimulation of hepatic gluconeogenesis and glycogenolysis, as well as elevated concentration of circulating free fatty acids as a result of increased adipocyte lipolysis [39]. All those mechanisms ultimately induce IR. However, the mechanism of catecholamine increase and the catecholamine levels are different in HFrEF and HFP EF. Catecholamine levels were reported to be higher in HFrEF compared to HFP EF. The increase in catecholamines may result from the activation of humoral neural stimuli secondary to the reduction in cardiac output in HFrEF and to alterations in left ventricular relaxation in HFP EF [11].

IR was reported to be an independent predictor of death among non-diabetic patients with HF [13], suggesting that impaired insulin sensitivity may play a role in the pathophysiology of HF progression. Some studies showed that the degree of IR correlated with the NYHA functional class of HF [34], the LVEF [13] and the peak oxygen consumption during a stress test [10, 13, 37]. In our study, when HFrEF and HFP EF patients were taken separately, participants with IR rather belonged to the highest NYHA functional classes and had higher NT-proBNP levels and lower LVEF compared to their counterparts without IR. This was observed after exclusion of the potential role of diabetes, overweight, hypertension or impairment of renal function, and whether or not patients had ischemic HF noting that ischemic HF was equally distributed in patients with or without IR. Furthermore, there was a trend for a more pronounced difference in HF severity according to the presence or absence of IR than according to the presence or absence of prediabetes (Tables 5 and 6). Thus, IR should also be considered for risk stratification in both HFrEF and HFP EF patients. Various mechanisms could explain how IR may alter cardiac function. In the presence of IR, the myocardium uses more free fatty acids and less glucose [45], and that increases heart vulnerability to ischemia and pressure load. Hyperinsulinemia increases sodium and fluid retention, enhances sympathetic nervous system activity [42] and may favor cardiac remodeling [46]. However, whether or not IR can be taken as a target for HF therapy remains uncertain. The Tayside observational study suggests that metformin may be beneficial in patients with HF and diabetes. This needs to be verified by a prospective clinical trial [47].

Finally, it is important to characterize glycemic and IR status in HF patients without previously known diabetes as it seems possible to reduce the incidence of diabetes in this population. A recent paper showed that compared with a strategy of general health education, a lifestyle intervention strategy can reverse glucose levels to normoglycemia in individuals with prediabetes [48]. In the HFrEF population included in the DAPA-HF trial, dapagliflozin treatment induced a 32% reduction in diabetes incidence and this effect was mainly driven by participants with prediabetes at baseline (HbA1c 5.7–6.4%) [9]. The potential benefit of sodium glucose cotransporter 2 (SGLT2)-inhibitors in diabetes prevention needs confirmation in trials of longer duration. It looks crucial as DAPA-HF participants who developed diabetes had a higher subsequent mortality than those who did not [9]. This additional benefit of SGLT2 inhibitors on diabetes prevention in HF patients could be reinforced in the presence of prediabetes and/or IR.

**Strengths and limitations**

Our study has major strengths. This was a prospective study including a reasonably large population of HF patients with either reduced or preserved left ventricle ejection fraction but a unique study design aiming to characterize glycemic and IR status in the total population. The population was relatively homogeneous due to the strict exclusion of patients with known or newly detected diabetes, hypertension or overweight and other factors that could have interfered with OGTT results or IR assessment. We were thus able to detect early stages of IR. The study has also some limitations. First, the study center was in Vietnam and the participants had a rather
low BMI, around 20 kg/m² in means: our findings may not be generalizable to other populations. In a previous study we showed that in non-overweight Vietnamese people, essential hypertension was associated with IR [49]. Overweight has been shown to be associated with a survival benefit in HF patients. Recent reports discussed a balanced reappraisal of this “obesity paradox” [50, 51]. Lean body weight in HF patients might result from weight loss associated with severe HF grade. However, our patients were in stable condition without evidence of recent weight loss. Future studies in non-overweight patients should take into account more accurately recent changes in body weight and carry out body composition evaluation. Second, the high prevalence of IR among our patients might be due to the high proportion of NYHA functional class III to IV (in 83% of the participants) and to the large predominance of ischemic HF. The association of IR with HF independently of the above-mentioned confounding factors needs to be confirmed in a population with other HF etiologies. Third, the cross-sectional design of the study prevents us from evaluating the impact of glycemic status and IR on HF-related outcomes. Fourth, HF patients were receiving different therapies, including renin angiotensin aldosterone system blockers, beta-blockers and diuretics, which may impact glucose metabolism. However, these treatments did not differ between patients with or without prediabetes or IR (data not shown).

Conclusion
The present data show that in Vietnam, in a population of participants with HF but no diabetes, overweight or hypertension, the prevalence of prediabetes and of IR is high in those with HFrEF as in those with HFrEF. Both prediabetes and IR are associated with more severe HF. This study contributes to better define the early stages of IR in HF patients with no metabolic disorders except mild hyperglycaemia and underlines the need to better assess all HF patients for their metabolic profile, including those without overweight. The study also provides arguments for HF as the culprit for IR. These interactions between HF, IR and prediabetes might carry prognostic and therapeutic implications. Intervention strategies should be proposed to HF patients with prediabetes aiming to reduce the risk of incident diabetes. Studies should be designed to test whether such strategies may translate into an improvement of further HF-related outcomes.

Abbreviations
BMI: Body mass index; EF: Ejection fraction; FIRI: Fasting insulin resistance index; HF: Heart failure; HFrEF: Heart failure with preserved left ventricle ejection fraction; HFrEF: Heart failure with reduced left ventricle ejection fraction; HOMA-IR: Homeostatic model assessment of insulin resistance; IFG: Impaired fasting glucose; IGT: Impaired glucose tolerance; IR: Insulin resistance; ISIT0: Fasting insulin sensitivity index; ISIT120: 2hH-insulin sensitivity index; NGT: Normal glucose tolerance; NT-proBNP: N-terminal pro Brain Natriuretic Peptide; NYHA: New-York Heart Association; OGTT: Oral glucose tolerance test; QUICKI: Quantitative insulin-sensitivity check index.

Acknowledgements
We are grateful to express our sincere gratitude to the Rectorate Board of Can Tho University of Medicine and Pharmacy and Hue University of Medicine and Pharmacy for creating favorable conditions for this study to be carried out. We would like to thank the Board of Directors of Can Tho Central General Hospital and the cardiologists who helped us to collect the samples, and especially thank the patients and their family members for taking their valuable time to participate in this study. Editorial assistance was provided by Sylvie Picard, Md, PhD, Dijon, France.

Author contributions
All authors have made substantial contributions. TKS and HVM conceived the study and designed the study protocol. NHT, HLTT, HAT and NHT organized, performed the study investigations and supported the recruitment of the patients. NHT and NT performed statistical analyses. TKS, NHT, HVM and PV wrote the first draft of the manuscript. All authors critically revised the manuscript for important intellectual content. All authors read and approved the final manuscript.

Funding
This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Availability of data and materials
The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
The study protocol was approved by our institution’s ethics committee, and informed consent for the procedure was obtained from each participant.

Consent for publication
Not applicable.

Competing interests
PV discloses the following potential conflicts of interest: lectures for Abbott, AstraZeneca, Bayer, Eli Lilly, Hikma Pharmaceuticals, Merck Sharp & Dohme, Novo Nordisk, Novartis, Pfizer, Sanofi; research grants from Abbott, Bristol-Myers Squibb–AstraZeneca, Novo-Nordisk; participation in expert committees for AstraZeneca, Boehringer Ingelheim, Novo Nordisk, Daiichi Sankyo, Sanofi, Servier. The other authors declare they have no conflict of interest.

Author details
1 Department of Internal Medicine, Can Tho University of Medicine and Pharmacy, Can Tho, Vietnam. 2 Science - Technology & External Relations Office, Can Tho University of Medicine and Pharmacy, Can Tho, Vietnam. 3 Cardiology Department, Can Tho Central General Hospital, Can Tho, Vietnam. 4 Department of Internal Medicine, Hue University of Medicine and Pharmacy, Hue University, Hue, Vietnam. 5 Unit of Endocrinology-Diabetology-Nutrition, Jean Verdier hospital, APHP, Sorbonne Paris Nord University, CINFO, CRNH-idF, Bondy, France.

Received: 3 February 2022   Accepted: 7 April 2022

Published online: 14 May 2022

References
1. Yancy CW, Jessup M, Bozkurt B, Butler J, Casey DE Jr, Colvin MM, et al. 2017 ACC/AHA/HFSA focused update of the 2013 ACCF/AHA Guideline for the Management of Heart Failure: a report of the American College of Cardiology/American Heart Association Task Force on Clinical
22. Otten J, Ahrén B, Olsson T. Surrogate measures of insulin sensitivity vs the
21. Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF, Turner RC.
20. Grundy SM, Cleeman JI, Daniels SR, Donato KA, Eckel RH, Franklin BA, et al.
19. Ponikowski P, Voors AA, Anker SD, Bueno H, Cleland JGF, Coats AJS, ESC
18. Groenewegen A, Rutten FH, Mosterd A, Hoes AW. Epidemiology of heart
17. Kenchaiah S, Evans JC, Levy D, Wilson PW, Benjamin EJ, Larson MG, et al.
16. Scherbakov N, Bauer M, Sandek A, Szabó T, Töpper A, Jankowska EA, et al.
15. Guglin M, Lynch K, Krischer J. Heart failure as a risk factor for diabetes
14. Stand E, Schnell O, McGuire DK. Heart failure considerations of antihyper-
13. Conrad N, Judge A, Tran J, Mohseni H, Hedgecott D, Crespillo AP, et al.
12. Heck PM, Dutka DP. Insulin resistance and heart failure. Curr Heart Fail
11. Swan JW, Anker SD, Walton C, Godsland IF, Clark AL, Leyva F, et al. Insulin
10. swallow the pill. Lancet. 2015;386:2545–50.
9. Dungan MH, Singh RM, Wise PH, Carter G, Alagha-band Zadeh J. A simple measure of insulin resistance. Lancet. 1995;346:120–1.
8. Cosentino F, Grant PJ, Aboyanos V, Bailey CJ, Cereillo A, Delgado V, ESC
7. Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF, Turner RC.
6. Grundy SM, Cleeman JI, Daniels SR, Donato KA, Eckel RH, Franklin BA, et al.
5. Dei Cas A, Khan SS, Butler J, Mentz RJ, Bonow RO, Cregg AP, et al. Impact of diabetes on epidemiology, treatment, and outcomes of patients with heart failure. JACC Heart Fail. 2015;3:136–45.
4. Cariello A, Cattano S, Cosentino F, Dombrowsky AC, Ithzik D, B&DV EASD Study Group, et al. Heart failure in type 2 diabetes: current perspectives on screening, diagnosis, and management. Cardiovasc Diabetol. 2021;20:218.
3. Tenenbaum A, Fisman EZ. Impaired glucose metabolism in patients with chronic heart failure: a possible etiologic link. J Am Coll Cardiol. 2009;54:109–16.
2. Nochioka K, Sakata Y, Miura M, Shiroto T, Takahashi J, Saga C, et al. Abnormal glucose tolerance contributes to the progression of chronic heart failure in patients with dilated cardiomyopathy. Hypertens Res. 2006;29:775–82.
1. Condon N, Judge A, Tran J, Mohseni H, Hedgecott D, Crespillo AP, et al. Temporal trends and patterns in heart failure incidence: a population-based study of 4 million individuals. Lancet. 2018;391:572–80.
42. Valensi P. Autonomic nervous system activity changes in patients with hypertension and overweight: role and therapeutic implications. Cardiovasc Diabetol. 2021;20:170.

43. Triposkiadis F, Karayannis G, Giamouzis G, Skoularigis J, Louridas G, Butler J. The sympathetic nervous system in heart failure physiology, pathophysiology, and clinical implications. J Am Coll Cardiol. 2009;54:1747–62.

44. Hartupee J, Mann DL. Neurohormonal activation in heart failure with reduced ejection fraction. Nat Rev Cardiol. 2017;14:30–8.

45. Witteles RM, Fowler MB. Insulin-resistant cardiomyopathy clinical evidence, mechanisms, and treatment options. J Am Coll Cardiol. 2008;51:93–102.

46. Sundström J, Lind L, Nyström N, Zethelius B, André R, Hales CN, Lithell HO. Left ventricular concentric remodeling rather than left ventricular hypertrophy is related to the insulin resistance syndrome in elderly men. Circulation. 2000;101:2595–600.

47. Evans JM, Doney AS, AlZadjali MA, Ogston SA, Petrie JR, Morris AD, et al. Effect of Metformin on mortality in patients with heart failure and type 2 diabetes mellitus. Am J Cardiol. 2010;106:1006–10.

48. Liu Y, Guo H, Wang Q, Chen J, Xuan Y, Xu J, Liu Y, Sun K, Gao Q, Sun Z, Wang B. Short-term effects of lifestyle intervention in the reversion to normoglycemia in people with prediabetes. Prim Care Diabetes. 2022;16:168–72.

49. Van Minh H, Thanh LC, Thi BN, do Trinh T, Tho TD, Valensi P. Insulinaemia and slight overweight: the case of Vietnamese hypertensives. Int J Obes Relat Metab Disord. 1997;21:897–902.

50. Gaddam KK, Ventura HO, Lavie CJ. Metabolic syndrome and heart failure—the risk, paradox, and treatment. Curr Hypertens Rep. 2011;13:142–8.

51. Mentz RJ, Kelly JP, von Lueder TG, Voors AA, Lam CS, Cowie MR, et al. Noncardiac comorbidities in heart failure with reduced versus preserved ejection fraction. J Am Coll Cardiol. 2014;64:2281–93.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Ready to submit your research? Choose BMC and benefit from:
• fast, convenient online submission
• thorough peer review by experienced researchers in your field
• rapid publication on acceptance
• support for research data, including large and complex data types
• gold Open Access which fosters wider collaboration and increased citations
• maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.
Learn more biomedcentral.com/submissions