**Gender differences in heart failure**

Renato Razzolini¹, Carlo Dal Lin¹

Summary. Heart failure is a growing health problem in the western world. Over the age of 40, the prevalence of heart failure is higher in men than in women, but this ratio is reversed after the age of 80. The aetiology of heart failure is mainly arterial hypertension in women, although ischemic heart disease can be important as well. The genetic basis for differences in aetiology is the chromosomal XY configuration in men, because several genes on the Y chromosome are associated with many cardiovascular risk factors. The pathophysiology of heart failure in women is mainly heart failure with preserved ejection fraction phenotype, probably because hypertrophy can easily take place without stretching the sarcomere. Symptoms are generally more severe in women. In particular, acute ischemic disease is often accompanied by left ventricular failure. Management should be similar to men, although in randomized studies women are poorly represented. It is said that ACE-inhibitors are less effective in women, but beta-blockers are more effective. Cardiac resynchronization therapy is certainly more effective in women than in men, while implantable cardioverter-defibrillators (ICD) are not, probably because sudden death is more uncommon in elderly women with heart failure of non-ischemic aetiology.

Key words. Heart failure, arterial hypertension, ischemic heart disease, ACE-inhibitors, beta-blockers.

Introduction

Heart failure is a growing health problem in the population of western world. This is the result of an increasing number of elderly individuals and their consequent long-term exposure to risk factors. Since life expectancy is approximately 5 years longer for women than for men, the prevalence of heart failure tends to be slightly higher in women. In this review we will briefly revise the gender-related features of heart failure, regarding epidemiology, pathophysiology, clinical presentation, treatment and prognosis.

Epidemiology

The prevalence of heart failure is shown in Table 1. After the age of 40, the prevalence of heart failure is higher among men at a ratio of (approximately) 2:1.

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>Men</th>
<th>Women</th>
</tr>
</thead>
<tbody>
<tr>
<td>20-39</td>
<td>0.2</td>
<td>0.4</td>
</tr>
<tr>
<td>40-59</td>
<td>1.5</td>
<td>0.7</td>
</tr>
<tr>
<td>60-79</td>
<td>7.8</td>
<td>4.5</td>
</tr>
<tr>
<td>80+</td>
<td>8.6</td>
<td>11.5</td>
</tr>
</tbody>
</table>

Table 1. Prevalence (%) of heart failure by gender and age (years). Modified from Taylor AL, 2015¹.

¹. Department of Cardiological, Thoracic and Vascular Sciences, Clinical Cardiology, University of Padua, Italy.

Invited paper received on 29 June 2015.
This condition is reverted after the age of 80. Since women outnumber men over the age of 80, the overall ratio of M:F in a given heart failure population is approximately 1:1.

The incidence of heart failure reduces over time in both sexes, but especially in women. The rate of hospital discharge is now convergent for men and women. This probably means that the rate of hospitalization is now the same for both sexes; previously this was not the case²-⁶.

**Genetic and physiological basis for gender difference**

The basic genetics of gender difference is the chromosome XY configuration for males and XX for females. Several genes on the Y chromosome are associated with many cardiovascular risk factors, namely increased blood pressure, increased low density lipoprotein cholesterol, and – in European males – the propensity to myocardial infarction.

In all ethnic groups, young men tend to have a higher mean systolic and diastolic pressure than young women. However, with advancing age the prevalence of hypertension is greater in women, meaning that the male:female ratio is reversed.

Chronic hypertension activates neurohormonal axis. Peripheral resistances are directly proportional to sympathetic drive in young men, but not in young women. In post-menopausal women, however, peripheral resistances increase with sympathetic drive, probably because oestrogen levels have declined. In actual fact, baroreflex sensitivity is modulated by oestrogen hormones. Hypertensive disorders of pregnancy increase the life-long risk for the development of hypertension or other cardiovascular pathologies later in life⁷-⁲¹.

**Pathophysiology**

The most important pathophysiological feature of heart failure in women is that left ventricular remodelling is oriented toward concentric hypertrophy. As a result, heart failure occurs in most cases with preserved ejection fraction (HFpEF). This is in contrast with heart failure in men, where the prevailing phenotype is heart failure with reduced ejection fraction (HFrEF).

The renin-angiotensin system would appear to be less activated. Consequently, fibrosis is less important at a myocardial level, although there is a general stiffening of the circulatory system with an increase in the effective afterload.

The reason for the preferential concentric hypertrophy and HFpEF instead of left ventricular dilatation and HFrEF is unknown. Some authors suggest that in normal conditions, myocyte diameter is reduced with respect to men, and it may therefore be possible to increase the amount of contractile proteins without stretching the sarcomere.

Response to acute ischemia is different, probably because of the presence of endogenous oestrogens. In short, ischemic damage is delayed, myocyte necrosis and even rupture are more difficult "in vitro", and late remodelling occurs less frequently. It should be noted that some of these experimental findings are inconsistent with well-known clinical knowledge. It is a well-known fact, for instance, that cardiac rupture after myocardial infarction occurs more frequently in women than in men, and that heart failure in acute coronary syndromes sets in commonly and early in women²⁸-⁴³.

**Clinical presentation**

As a rule of thumb, symptoms are worse in women than in men at a given ejection fraction and/or isch-
emic burden. As far as ejection fraction is concerned, this is easily explained: reduced ejection fraction is not the main cause of heart failure in women. On the contrary, the main causes are increased left ventricular diastolic dysfunction and ventriculo-arterial coupling, due to increased systolic and arterial elastance (Figure 1). Therefore, a similarly increased left ventricular end diastolic pressure – the direct cause of respiratory symptoms in heart failure – is reached with a more depressed ejection fraction in men respect to women.

The increased incidence of heart failure early in acute coronary syndromes may have the same explanation. Failure by the myofiber to relax is an early mechanical manifestation of acute ischemia. Clearly, if a diastolic dysfunction is already present, symptoms appear earlier.

Although women suffer more severe symptoms, they tend to have a longer survival, which is not uniformly acknowledged. However, morbidity is much higher in women, who therefore experience a generally worse quality of life. It is unclear whether more severe symptoms entail an increased rate of hospitalization. Indeed, this may not be the case, because most symptoms are non-specific and are often – at least partly – overlooked.

One interesting feature is that depressive symptoms often accompany heart failure in women. In a community sample of 338 non-institutionalized adults aged 45 years old, depressive symptoms were measured using the Beck depression inventory. Women scored significantly higher than men. Adjusted score increased linearly with heart failure stages A to C in women, while in men only stage C was associated with a significantly higher score. It should be noted that depression worsens prognosis.

**Treatment**

Treatment of heart failure is not gender-specific. Therefore, as a general rule in selecting therapeutic strategies current guidelines should be followed. That said, one cannot help noting that in clinical trials less than 30% of participants are female. Considerations about gender-related activity of some drugs or drug classes derive from post-hoc analyses.

**General consideration**

The percentage of lean mass is lower in females; therefore, volume distribution of liposoluble drugs is greater, whereas volume distribution of hydrosoluble substances is smaller.

Cytochrome P<sub>450</sub> isoenzyme activity differs in men and women (Table 2). The consequences are particularly important for beta-blockers and calcium blocking agents, whose action is respectively increased and decreased in females.
Drug classes

Ace-inhibitors

Although there is no substantial data, current opinion favours the idea that ACE-inhibitors are less effective in women than in men. In Consensus-I and SOLVD-treatment trials, mortality reduction was 5% in women as compared with 30-40% in men. A cough is certainly a more frequent, disturbing side effect in women. One reason for the diminished efficacy of ACE-inhibitors could be the prevalence of HFpEF in females, where RAAS blockage is less useful. It is interesting to note that this gender-related difference does not appear to be present in angiotensin-II receptor blockers. In short, ACE-inhibitors would appear to be partially effective in the general female population with heart failure, and particularly effective in symptomatic women with reduced ejection fraction.

Beta blockers

Beta blockers are certainly useful in women, with some slight differences seen between the various molecules. In some cases, their effectiveness would appear to be higher than in men, but this may be a false impression, because natural history of heart failure is more favourable in women. The relative efficacy is the same. The most probable explanation is that the cytochrome P450 isoenzyme involved in metabolism of betablockers has an enhanced action in men, so that drug activity in women is enhanced.

Device therapy

Device treatment has significantly improved the outlook of heart failure patients. Indeed, ICD treatment, both in primary and secondary prevention, avoiding sudden death, has dramatically improved their survival. Once again, gender-specific data is lacking, but the impression is that ICD is not as useful in women as it is in men, probably because sudden death is less frequent in women with heart failure. A possible exception exists in the form of ischemic heart disease, where incidence of sudden death is appreciable.

By contrast, cardiac resynchronization therapy (CRT) is particularly effective in women. Reverse remodelling is easily induced, measurable as reduced left ventricular end systolic volume, and a survival advantage therefore ensues. A recent meta-analysis considers that the relative probability to reduce left ventricular end systolic volume ≥15% is 1.12 (0.99-1.26) in women with respect to men. One possible explanation is that normal QRS duration is slightly shorter in females than in men, and therefore, at a given QRS lengthening, the mechanical damage may be more severe.

Conclusions

There are certainly differences in pathophysiology, aetiology and clinical presentation of heart failure in men and women. These differences should probably prompt different treatment strategies. However, we do not presently have sufficient data to suggest gender-specific therapies. General guidelines should be followed, whilst awaiting gender-targeted studies on heart failure.

Key messages

- Later-in-life heart failure is more prevalent in women than in men.
- Aetiology is especially hypertension, although ischemic heart disease and diabetes play an important role.
- The most common phenotype is heart failure with preserved ejection fraction.
- Treatment should follow general guidelines, although in randomized studies the female gender is poorly represented.
- Cardiac resynchronization therapy is particularly efficient.
References


Ital J Gender-Specific Med 2015; 1: 15-20

44. Scantlebury DC, Borlaug BA. Why are women more likely than men to develop heart failure with preserved ejection fraction? Curr Op Cardiol 2011; 26: 562-8.

Correspondence to:
Renato Razzolini
Department of Cardiological, Thoracic and Vascular Sciences, Clinical Cardiology
University of Padua, Italy
email renato.razzolini@unipd.it