Underlying hemodynamic differences are associated with responses to tilt testing

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Aim of this study was to explore whether differences in resting hemodynamic parameters may be associated with tilt test results in unexplained syncope. We analyzed age, gender, systolic (SBP), diastolic blood pressure (DBP) and heart rate (HR) by merging three large databases of patients considered likely to be of vasovagal reflex etiology, comparing patients who had tilt-induced reflex response with those who did not. Tilt-induced reflex response was defined as spontaneous symptom reproduction with characteristic hypotension and bradycardia. Relationship of demographics and baseline supine BP to tilt-test were assessed using logistic regression models. Individual records of 5236 patients (45% males; mean age: 60 ± 22 years; 32% prescribed antihypertensive therapy) were analyzed. Tilt-positive (n = 3129, 60%) vs tilt-negative patients had lower SBP (127.2 ± 17.9 vs 129.7 ± 18.0 mmHg, p < 0.001), DBP (76.2 ± 11.5 vs 77.7 ± 11.7 mmHg, p < 0.001) and HR (68.0 ± 11.5 vs 70.5 ± 12.5 bpm, p < 0.001). In multivariable analyses, tilt-test positivity was independently associated with younger age (Odds ratio (OR) per 10 years: 1.04; 95% confidence interval (CI), 1.01–1.07, p = 0.014), SBP ≤ 128 mmHg (OR: 1.27; 95%CI, 1.11–1.44, p < 0.001), HR ≤ 69 bpm (OR: 1.32; 95%CI, 1.17–1.50, p < 0.001), and absence of hypertension (OR: 1.58; 95%CI, 1.38–1.81, p < 0.001). In conclusion, among patients with suspected reflex syncope, younger age, lower blood pressure and lower heart rate are associated with positive tilt-test result.

Tilt-testing (TT) was first introduced for clinical diagnosis of reflex syncope in 19861, and has over the past decades been criticized for failing to provide discrimination between reflex (vasovagal) and other more serious causes of syncope, such as arrhythmic syncope, despite its acceptable sensitivity and specificity in true positive and negative subjects2. Thus, it is claimed that TT offers limited diagnostic value in those for whom it is most needed5. In this study we aimed to offer an alternative explanation for why some subjects with likely reflex syncope faint during tilt testing and others do not. The ISSUE 3 (International study of syncope of unknown etiology) set out to identify patients with recurrent syncope, unexplained but thought likely to be of reflex origin in older patients who had received a dual-chamber rate-drop response pacemaker after documentation of an asystolic reflex episode4. Unexpectedly, patients with positive pre-implant-tilt were relatively unprotected against recurrent syncope during the pacing phase. In contrast, patients with negative pre-implant tilt-testing had a very low incidence of recurrent syncope5. The new interpretation of tilt-testing incorporated a susceptibility to vertical posture stress, possibly co-existing with any cause of syncope. This so-called ‘hypotensive susceptibility’ is caused by a vertically induced fluid shift, and is associated with the hypotensive (vasodepressor) aspect of reflex syncope3.

Recently, we have identified a specific cardiovascular profile in patients with reflex syncope, characterized by lower systolic blood pressure (SBP), elevated heart rate (HR), and diastolic (DBP) compared with the general population6. Other recent studies have emphasized the primary role of hemodynamic changes in the mechanism...
Methods

Study population and design. We analyzed individual values of resting supine BP and HR collected between 2003 and 2019 in three large databases of patients who had undergone TT for unexplained syncope, in tertiary syncope investigation units of hospitals in Lavagna, Italy (n = 2798), Florence, Italy (n = 805), and Malmö, Sweden (n = 1633). Consecutive patients who had undergone TT for diagnosis of syncope were included. Those with classical orthostatic hypotension (OH) and suspected postural orthostatic tachycardia syndrome (POTS) were excluded. Hypertension was defined as a clinical diagnosis of hypertension and antihypertensive medication reported by the patient. Despite the long study period, indications, methodology and interpretation of TT results remained unchanged during this time and were very similar to the recommendations of the current ESC syncope guidelines.

TT was performed according to the “Italian protocol” and endorsed by the most recent ESC guidelines on Syncope. The protocol consisted of at least 10-min supine period, a 20-min passive phase at a tilt angle of 70°, followed by a 15-min nitroglycerine-potentiated phase (300–400 mcg administered sublingually) if syncope was not induced during the passive phase. Positive response was defined as reproduction of spontaneous symptoms in the presence of bradycardia and hypotension. In all three centers, baseline hemodynamic data were obtained in the supine position prior to TT, using validated non-invasive beat-to-beat hemodynamic monitors, Task Force Monitor (CNSystems Medizintechnik GmbH, Graz, Austria) in Florence and Lavagna, and Nexfin (BMEYE, Amsterdam, Netherlands) or Finapres Nova monitors (Finapres Medical Systems, PH Enschede, Netherlands) in Malmö. The monitors were calibrated before measurement using brachial cuff and oscillometric method according to the manufacturer’s instructions. As the monitors render beat-to-beat data, an average value of a hemodynamically stable period of 10 (Florence and Lavagna) or 30 s (Malmö) was recorded in the database. Data on baseline heart rate were unavailable in one of the three study-centers (Florence, Italy). The patient information was de-identified before merging databases. Ethical approval for original data collection was obtained by an appropriate institutional review board: “Comitato Etico Regionale” of Regione Liguria, Italy for Lavagna cohort, “Comitato Etico Area Vasta Centro, AOU Careggi, Florence”, for Florence cohort, and Regional Ethical Review Board in Lund, Sweden for Malmö cohort, and all participants gave their written informed consent. According to Italian and Swedish law, the use of anonymous patient data previously collected for patient care, as was the case here, did not require additional ethical approval.

Statistical analyses. We retrieved the following data: age, gender, resting SBP and DBP, HR, and use of antihypertensive drugs although HR were not available in the Florence database. All the datasets were subsequently merged into one study population. Additionally, rate-pressure product (HR x SBP), a marker of cardiovascular fitness and myocardial oxygen consumption, was calculated for merged Lavagna and Malmö databases.

Continuous data are shown as mean ± standard deviation, whereas frequencies are used to describe categorical data. The method of Kolmogorov and Smirnov was used to check the normality of distributions. Continuous variables were compared by means of the paired Student’s t-test. Paired and multiple proportions were compared by means of Pearson’s chi-square test. Stepwise multiple regression analysis was used to identify the independent factors associated with TT positivity. Multivariable logistic regression was adjusted for age, gender, SBP, and HR, whereas presence of hypertension was adjusted for age, gender and HR only. In addition, the results of the three centers were separately analyzed.

Analyses were performed using the Statistical Analysis System Software (version 9.4; SAS Institute, Cary, NC, USA; https://www.sas.com/sv_se/home.html) and IBM SPSS Statistics software (version 26.0; SPSS Inc., Chicago, IL, USA; https://www.ibm.com/products/spss-statistics). Statistical significance was set at the 0.05 level and all p-values were two-sided.

Results

Study population. The total study population consisted of 5236 patients; 55% of whom were females, the mean age was 60 ± 22 years. A total of 1655 patients (32%) were prescribed antihypertensive therapy. TT induced reflex syncope in 3129 (60%) patients and did not induce reflex syncope in 2,107 (40%) patients. Five-hundred-and sixty tests were positive during passive TT, and 2569 during drug-potentiated phase. TT was positive in 61% of males and 59% of females.

Comparison between patients with and without tilt-induced reflex response. Compared with TT-negative patients, TT-positive patients had lower baseline SBP (127.2 ± 17.9 vs 129.7 ± 18.4, p < 0.001), DBP (76.2 ± 11.5 vs 77.3 ± 11.7, p < 0.001) and HR (68.0 ± 11.5 vs 70.5 ± 12.5, p < 0.001). These differences were present both in males and females with small, but statistically significant, differences: baseline SBP was similar in males and females, but males had higher DBP whereas females had higher HR than males (p = 0.001 for DBP and HR) (Fig. 1).

Comparison between subgroups. In univariate analyses (Table 1), we found a significantly higher rate of TT-induced reflex response among syncope patients below age 30 years, and in patients without history of hypertension. Lower baseline SBP below the mean value of 128 mmHg and lower baseline HR below the mean value of 69 bpm was also significantly associated with TT-induced reflex response. In a multivariable-adjusted
Figure 1. Gender-stratified systolic blood pressure, diastolic blood pressure and heart rate in tilt-positive and tilt-negative patients. DBP diastolic blood pressure; HR heart rate; SBP systolic blood pressure; TT− tilt-negative; TT+ tilt-positive. Resting systolic blood pressure, diastolic blood pressure and heart rate among tilt-positive and tilt-negative patients stratified by gender who were investigated using Italian tilt-test protocol for suspected syncope. Males, tilt-positive, n = 1428, tilt-negative, n = 923; females, tilt-positive, n = 1701, tilt-negative, n = 2107. All the values are shown as mean ± 1 SE. The bars show the ± 95% confidence limit (2 SEs).

Table 1. Univariable and multivariable analysis of factors predicting tilt test positivity in patients investigated for unexplained syncope using the Italian tilt test protocol. SBP systolic blood pressure, HR heart rate.

*Adjusted for age, gender and heart rate.

| Variables | Total syncope population (n = 5236) | Tilt-positive patients (n = 3129) | Univariable analysis p value | Multivariable analysis p value |
|-----------|-------------------------------------|----------------------------------|-----------------------------|-------------------------------|
| Age subgroups |                                      |                                  |                             |                              |
| 10–29 years | 748                                 | 493 (66)                         | 0.006                       | 0.016                         |
| 30–59 years | 1415                                | 848 (60)                         |                             |                              |
| ≥ 60 years  | 3073                                | 1788 (58)                        |                             |                              |
| Gender |                                      |                                  |                             |                              |
| Males | 2351                                 | 1428 (61)                         | 0.20                        | 0.10                          |
| Females | 2885                                 | 1701 (59)                         |                             |                              |
| Baseline SBP at the time of tilt testing |                                      |                                  |                             |                              |
| > 128 mmHg | 2443                                | 1358 (56)                         | 0.0001                      | 0.0001                        |
| ≤ 128 mmHg | 2793                                | 1771 (63)                         |                             |                              |
| Baseline HR at the time of tilt testing |                                      |                                  |                             |                              |
| > 69 bpm | 2088                                 | 1176 (56)                         | 0.0001                      | 0.0001                        |
| ≤ 69 bpm | 2324                                 | 1490 (64)                         |                             |                              |
| Hypertension* |                                      |                                  |                             |                              |
| Yes | 1655                                 | 864 (52)                          | 0.0001                      | 0.0001                        |
| No | 3581                                 | 2265 (63)                         |                             |                              |

Abbreviations: DBP, diastolic blood pressure; HR, heart rate; SBP, systolic blood pressure; TT−, tilt-negative; TT+, tilt-positive.
 logistic regression model (Table 1), younger age, lower BP and HR, and absence of hypertension remained the independent correlates of test positivity. In particular, a TT-induced reflex response was independently associated with younger age (Odds Ratio (OR) [per 10 years]: 1.04; 95% confidence interval (CI), 1.01–1.07, p = 0.007), SBP below 128 mmHg (OR: 1.27; 95%CI 1.11–1.44, p < 0.001), and HR below 69 bpm (OR:1.32; 95%CI 1.17–1.50, p < 0.001). Absence of hypertension diagnosis increased the probability of a positive TT-induced reflex response by over 50% (OR: 1.58; 95%CI 1.38–1.81, p < 0.001).

Exclusion of one database did not substantially affect the overall results except for the impact of low SBP on TT positivity, which was attenuated after exclusion of Lavagna data in the multivariable-adjusted (but not univariate) logistic regression model (p = 0.083). Tilt testing results and resting hemodynamic parameters stratified according to the study center are shown in Table S1. The associations of BP with tilt positivity were weaker in Florence cohort, being significant for DBP only. The rate-pressure product was significantly lower among TT-positive patients across age and sex strata with exception of youngest females < 30 years (Figure S1). All rate-pressure product values were within the normal range (< 12,000)13.

Discussion

In this three-center analysis of syncope patients with high pre-test probability of reflex mechanism, we found a slightly different hemodynamic pattern among those who were tilt-test positive. Males and females who had induction of reflex syncope during TT had lower baseline SBP, DBP, and HR compared with those who did not. We propose that patients more resistant to orthostatic stress induced by TT, who have slightly higher BP and HR, may have greater hemodynamic reserve in the face of gravitational challenge.

Possible interpretation of tilt test results. Our results suggest that patients with TT-induced reflex syncope may have increased resting vagal tone and/or lower sympathetic tone, both expressed by lower HR, and narrower hemodynamic margins in the face of orthostatic stress, expressed by lower initial BP. Hypotensive susceptibility is suspected to exist in most people, particularly in individuals with a history of syncope7. Its identification is thought to play a key role in the management of syncope and selection of effective therapy, such as guiding pacemaker therapy in reflex syncope18, as well as to be the reason for the good response to pacing in tilt-negative patients7. In writing of the European Society of Cardiology Guidelines on Syncope 2018, the concept of hypotensive susceptibility was further extended to anticipate that some patients with reflex syncope were constitutionally predisposed to reflex syncope based on presenting a low blood pressure—a low blood pressure phenotype9.

We have previously reported that individuals investigated due to unexplained syncope of likely reflex mechanism have in general lower SBP, but higher HR and DBP compared with normal population6. According to this study, patients who reproduce reflex syncope during TT have lower not only SBP, but also DBP and HR, and may thus represent a more vulnerable subset of syncope population. Recent studies have indicated that TT positivity is associated with neuroendocrine activation characterized by excess epinephrine and vasopressin release15,16, whereas adrenomedullin seems to play a protective role, probably reducing vessel permeability and acting against intravascular volume reduction during orthostasis15. Further, older age and higher resting SBP have been suggested to predict lower susceptibility to positive tilt-testing15,17. These findings were supported by Lindenberger et al. who observed upregulation of vasopressin release, reduced cardiac filling and cardiac output in women prone to reflex syncope16. It might be interpreted that vasopressin upregulation is a secondary effect of a relative hypovolemia, compared with non-vasovagal subjects, suggesting a key role of discrete water and electrolyte imbalance in increased tendency to syncope. There, also, remains a tenable possibility that the documented endocrine changes are precipitated by the developing adverse hemodynamic picture i.e. reduced venous return and cardiac filling. Combination of lower circulating blood volume and excessive blood pooling in capacitance vessels during orthostatic challenge may abruptly reduce stroke volume during TT, with a compensatory HR increase, as shown by Buszko et al.18.

Those with a more pronounced TT sensitivity demonstrate increased epinephrine and vasopressin release during tilt testing and thus appear to be in greater need of circulatory compensation as their resting SBP and DBP are lower15,17. This may provoke response from the central nervous system, leading to reflex syncope in extreme situations20,21. As previously shown, this response occurs when cerebral tissue oxygenation is strongly compromised22.

Although female gender, especially at younger age, is associated with higher syncope incidence23, only younger age but not gender was independently predictive of TT positivity. The protective factors against TT intolerance were resting SBP above 128 mmHg, heart rate above 69 bpm and presence of hypertension, even while prescribed antihypertensive treatment.

Higher HR might indicate less vagal tone and more efficient chronotropic compensatory mechanisms. As SBP mainly increases with greater stroke volume, reduced arterial elastance, and, to a lesser degree, with elevated peripheral vascular resistance23, higher SBP may suggest that those who are capable of tolerating TT have greater intravascular volume, better cardiac filling, and higher systemic vascular resistance compared with those who developed vasovagal syncope during TT. Interestingly, the association of combined heart rate and systolic blood pressure with TT positivity was markedly attenuated in younger women, who are genetically most predisposed to reflex syncope22, a fact that deserves more studies. Hypertension, on the other hand, implies chronic sympathetic activation with increased total peripheral resistance and elevated arterial tone in the precapillary vascular bed26. Arterial hypertension is detrimental for long-term cardiovascular integrity and promotes end-organ damage27, but hypertensive patients seem to be more resistant to orthostatic stress, either due to altered hemodynamic reserve, increased circulating blood volume, chronic neuroendocrine activation and arterial vasoconstriction or by baroreceptor resetting.
The differences in observed hemodynamic parameters between tilt-positive and tilt-negative patients were small, yet highly statistically significant. Despite these differences having limited clinical applications, they are important for clarifying pathophysiological mechanisms behind reflex syncope susceptibility in some individuals. Future studies might broaden the concept of hemodynamic prerequisite for reflex syncope by using Model-based estimation of cardiac stroke volume and by applying prolonged and more accurate BP and HR assessment methodology such as 24-h ambulatory BP monitoring and heart-rate variability, as well as peripheral vascular resistance, blood volume, and venous return estimation.

**Strengths and limitations.** The strengths of this three-center study lie in the large number of patients included and almost identical examination protocol, thus minimizing potential inter-center variability. There are some limitations that should be mentioned. Data on baseline heart rate were unavailable in one of the study centers (Florence, Italy), however, despite this limitation, the results in the remaining two study-centers demonstrated consistency. The study is retrospective with all typical limitations for retrospective analyses. Lower overall stress and sympathetic drive could have contributed to the lower SBP, DBP and HR observed in patients with TT-induced reflex syncope. We are unable to confirm or refute this possibility, however, this does not contradict our study hypothesis of a hemodynamic phenotype contributing to reflex syncope susceptibility. Further, patients without induction of reflex syncope during TT may have suffered syncope due to competing mechanisms such as cardiac arrhythmia but the final diagnoses in this patient group were not available. Finally, in believing that the constitutional nature of a low blood pressure phenotype contributes importantly to hypotensive susceptibility, we found it necessary to include a wide range of ages and hypertensive patients, some of whom were likely to have a similar susceptibility prior to developing their hypertensive disease. Our clinical observations suggest also that some patients reporting a history of reflex syncope in youth that resolves in mid-life, sustain recurrence to have a similar susceptibility prior to developing their hypertensive disease. Our clinical observations suggest also that some patients reporting a history of reflex syncope in youth that resolves in mid-life, sustain recurrence.

**Conclusions**

Patients who develop reflex syncope during tilt testing have significantly lower blood pressure and heart rate compared with tilt-negative patients. In contrast, advanced age and hypertension are two important factors diminishing tilt-testing positivity. Our findings support the concept of a low heart rate and low blood pressure phenotype contributing to reflex syncope susceptibility during orthostatic stress.

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**Author contributions**

A.F., G.R., R.S., and M.B. wrote the main manuscript text and M.B. prepared Fig. 1 and Figure S1. A.F., G.R., P.T., M.J., M.R., I.M., A.C., N.C., V.H., A.U., M.B., G.P. provided the data for the manuscript. P.T., M.J., M.R., I.M., A.C., N.C., V.H., A.U., B.O., and G.P. reviewed the manuscript before submission. A.F. and M.B. prepared the final manuscript.

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**Competing interests**

The authors declare no competing interests.

**Additional information**

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