Review

Syncope and the risk of sudden cardiac death: Evaluation, management, and prevention

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Abstract

Syncope is a clinical syndrome defined as a relatively brief self-limited transient loss of consciousness (TLOC) caused by a period of inadequate cerebral nutrient flow. Most often the trigger is an abrupt drop of systemic blood pressure. True syncope must be distinguished from other common non-syncope conditions in which real or apparent TLOC may occur such as seizures, concussions, or accidental falls. The causes of syncope are diverse, but in most instances, are relatively benign (e.g., reflex and orthostatic faints) with the main risks being accidents and/or injury. However, in some instances, syncope may be due to more worrisome conditions (particularly those associated with cardiac structural disease or channelopathies); in such circumstances, syncope may be an indicator of increased morbidity and mortality risk, including sudden cardiac death (SCD). Establishing an accurate basis for the etiology of syncope is crucial in order to initiate effective therapy. In this review, we focus primarily on the causes of syncope that are associated with increased SCD risk (i.e., sudden arrhythmic cardiac death), and the management of these patients. In addition, we discuss the limitations of our understanding of SCD in relation to syncope, and propose future studies that may ultimately address how to improve outcomes of syncope patients and reduce SCD risk.

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1. Introduction

Syncope is a clinical syndrome defined as a relatively brief and self-limited transient loss of consciousness (TLOC) caused by a period of inadequate cerebral nutrient flow. Most often the trigger is an abrupt drop of systemic blood pressure. True syncope must be distinguished from other common non-syncope conditions in which real TLOC may have occurred such as seizures or concussions, or in which TLOC may seem to have occurred such as with accidental falls or psychogenic pseudosyncope.

The causes of syncope are diverse, but in most cases the cause itself is relatively benign (e.g., reflex or orthostatic faints) with the main risks being the consequences of loss of postural tone, such as falls, leading to accidents and injury. However, in some instances, syncope may be due to more worrisome conditions (particularly those associated with cardiac structural disease or channelopathies), and, in such circumstances, syncope may be an indicator of increased morbidity and mortality risk, including sudden cardiac death (SCD). Further, the likelihood of a cardiac origin for syncope, and the consequent increased risk of serious adverse events, are greater in older patients than in the young, paralleling the inevitable development of underlying serious structural heart disease with advancing age.

Overall, morbidity and mortality in syncope patients is low, but 1-year mortality can reach 33% in certain subgroups of patients having a cardiac etiology of syncope. Consequently, establishing an accurate diagnosis and instituting effective preventive measure is essential [1]. Unfortunately, however, rates of unexplained syncope remain high, emphasizing the importance of both developing more effective diagnostic strategies and promoting their acceptance by clinicians [2].

2. Sudden cardiac death: definition, etiology, and pathophysiology

SCD is a term used to refer to a mode of cardiac death, and is frequently used as an outcome of interest in research and epidemiological studies. While there has been some debate, the definition of SCD by Myerburg and Castellanos is widely accepted: “A natural death due to cardiac causes, heralded by abrupt loss of consciousness, within 1 hour after the onset of acute symptoms or an unwitnessed, unexpected death of someone seen in a stable medical condition less than 24 hours previously with no evidence of a non-cardiac cause.” [3,4]. In these cases, it is assumed that there is a sudden cessation of organized cardiac electrical activity, leading to hemodynamic collapse. If circulation is restored, either spontaneously or through intervention (e.g., defibrillation, antiarrhythmic drugs), the event is referred to as either a sudden cardiac arrest (SCA) or SCD, but, if fatality occurs, it can only be termed as SCD. If, on the other hand, resuscitation is initially effective, but the patient dies somewhat later, then it is classified as a ‘non-sudden’ cardiac death.

Abnormal electrical activity associated with SCD may be broadly classified as being tachyarrhythmias and non-tachyarrhythmias. The former includes ventricular fibrillation (VF) and pulseless ventricular tachycardia (VT), which most often arise from cardiac causes; only rarely are they due to non-cardiac causes (e.g., pulmonary embolism). In any case, these life-threatening arrhythmias require urgent therapy, specifically direct current (DC) shock and often other interventional cardiac procedures [5,6]. The non-tachyarrhythmia group includes pulseless electrical activity (PEA), asystole, and extreme bradycardia. While commonly associated with non-cardiac factors (e.g., major pulmonary embolism), PEA may have a primary cardiac cause, including severe pump failure and acute coronary syndrome. Secondary cardiac causes of PEA include those that occur following spontaneous or electrical termination of VT or VF [7].

In industrialized countries and, most likely as well, in developing economies, the most common cardiac cause of SCD is myocardial ischemia due to atherosclerotic coronary artery disease (CAD) [4,8]. However, SCD may also be a complication of a wide variety of other cardiac conditions [Table 1] [4,9]. Finally, there are non-arhythmic forms of SCD, such as aortic dissection, massive pulmonary embolism, cardiac tamponade, and atrial myxoma. In this report, we use “cardiac syncope” to refer to syncope secondary to a cardiac arrhythmia. Non-arhythmic forms are reasonably categorized separately as of ‘structural cardiovascular origin’.

3. Syncope and its prognosis

The mortality associated with syncope, including SCD risk, is greatest in those cases in which syncope is of a cardiac cause. Mortality rates of 18% to 30% at 1 year, as compared to only 6% in adult patients with syncope of unknown origin (the majority of which are probably reflex or orthostatic), have been reported [10,11]. Pre-syncope, in at least one study, has been shown to be as important as true syncope from a prognostic perspective, and, therefore, is similarly managed [12].

The report by Soteriades et al., derived from Framingham Heart Study data, despite being limited by incomplete diagnostic testing and the uncertain ‘neurological syncope’ category, was among the first to highlight the importance of ‘cardiac cause’ as a major mortality risk factor in syncope patients [11]. Of community-dwelling participants in Framingham, the incidence of a first report of syncope was 6.2 per 1000 person-years. Of those, 21.2% were believed to be vasovagal syncope, 9.5% cardiac syncope, 9.4% orthostatic, and 36.6% were of unknown cause. Multivariable-adjusted hazard ratios among individuals with presumed cardiac syncope were 2.01 (95%CI 1.48 to 2.73) for death from any cause, 2.66 (95%CI 1.69–4.19) for myocardial infarction or death from coronary heart disease (including sudden and non-sudden death), and 2.01 (95%CI 1.06 to 3.80) for fatal or nonfatal stroke. There was no increased risk of cardiovascular morbidity or mortality associated with a presumptive diagnosis of vasovagal syncope. In brief, although the Framingham report did not assess SCD risk per se,
and, in the case of young athletes, unexpected deaths have become terms of mortality risk; however, unexpected deaths may occur, increased mortality risk.

It showed a concerning association between cardiac syncope and are summarized in Table 2. However, an important caveat with respect to the occurrence of certain channelopathies was not yet fully appreciated. Age is an important factor. In this regard, Del Rosso et al. compared the diagnosis of the cause of syncope was possible by history alone other than coronary atherosclerosis.

Table 1
SCD causes and contributing factors other than coronary atherosclerosis.

| Bradyarrhythmias (e.g. complete heart block) | Cardiomyopathies |
|--------------------------------------------|------------------|
| - Alcohol                                  | - Dilated – idiopathic |
| - Chagas disease                           | - Hereditary      |
| - Hypertrophic                             | - Hypertensive    |
| - Infertile (e.g., amyloid, sarcoid, etc)  | - Peripartum      |
| - Cellular conduction disturbances (e.g., ARVC) | - Takotsubo (Stress-induced) |
| - Brugada syndrome                         | Channelopathies   |
| - Catecholaminergic polymorphic VT         |                  |
| - Early repolarization                      |                  |
| - Long and Short QT syndrome               |                  |
| Conduction system abnormalities (Wolf-Parkinson-White) |                  |
| Congenital coronary-artery anomalies       |                  |
| Congenital heart diseases                  |                  |
| Coronary artery abnormalities (e.g., spasm, dissection, embolism) |                  |
| Laminopathies                              |                  |
| Left ventricular noncompaction             |                  |
| Mechanical interference of venous return (e.g., pulmonary embolism, tamponade) |                  |
| Neuromuscular diseases (e.g., muscular and myotonic dystrophies) |                  |
| Pulmonary hypertension (primary and secondary causes) |                  |
| Myocardiitis                               |                  |
| Valvular disease (mitral valve prolapse, aortic stenosis) |                  |

ARVC represents arrhythmogenic right ventricular cardiomyopathy; SCD, sudden cardiac death; VT, ventricular tachycardia

4. Patient evaluation

As a rule, when patients present for evaluation of a presumed TLOC event, they use non-specific terms to describe their symptom experience; common descriptors in North America are ‘collapse’ or ‘blackouts’ or falls. However, the clinician should not assume that these episodes were ‘true syncope,’ because that might lead to overlooking other potential causes of real or apparent TLOC (e.g., seizures, accidents, drug abuse, psychogenic pseudosyncope/seizure). Consequently, the first hurdle faced by the clinician is determining whether the episode(s) was due to syncope, or of some other cause of real or apparent TLOC. For purposes of this communication, we assume that the basis for collapse was ‘syncope’ as defined earlier, and consequently the next hurdle is ascertaining the underlying cause.

4.1. Medical history

The foundation for defining the etiologic basis for syncope is a comprehensive medical history taken by an experienced clinician. Importantly, a detailed account should be obtained from the patient and, when possible, witnesses. The initial assessment should include careful documentation of several symptomatic episodes, looking for similarities suggesting a causal diagnosis. Finally, it is crucial to document pre-existing medical conditions, ongoing and newly introduced drug therapy, and family history.

In general, a thorough detailed clinical history taken by an experienced practitioner will be sufficient to differentiate syncope from non-syncope in most cases, and even provide a reasonable explanation for the collapse in 40 to 70% of cases. However, in many instances distinguishing true syncope from non-syncope collapse may be challenging and/or a plausible etiologic diagnosis may not be evident.

Once it is clear that syncope has occurred, differentiating benign causes from those that could be life-threatening is essential. From a mortality and SCD perspective, it is essential to identify ‘cardiac’ causes of syncope (i.e., cardiac structural causes as well as channelopathies), because these comprise the highest mortality risk cases.

Certain clinical symptoms and features have been associated with cardiac syncope and are summarized in Table 2. Several patient characteristics (e.g., age or the presence of structural heart disease) drastically increase the pre-test probability of cardiac syncope. For example, Alboni et al. found that underlying heart disease was an independent predictor of cases of syncope, and the absence of heart disease excluded a cardiac cause of syncope in 97% of patients. However, an important caveat with respect to the latter findings is that, at the time of the study, the importance of certain channelopathies was not yet fully appreciated. Age is another important factor. In this regard, Del Rosso et al. compared findings in patients above and below age 65; they reported that the diagnosis of the cause of syncope was possible by history alone in 26% of younger and only 5% of older patients. Finally, the value of medical history often becomes less reliable as patients age. For instance, it is now recognized that older patients with syncope may have a period of retrograde amnesia that undermines their recollection of preceding events. In fact, they often insist that they never lost consciousness. Only with the aid of reliable witnesses can the story become clear.
Table 2
Common clinical predictors in syncope subtypes.

| Clinical features that suggest a diagnosis on initial evaluation |
|---------------------------------------------------------------|
| **Neurally mediated syncope:** |
| - Absence of heart disease |
| - Absence of trauma |
| - After exertion |
| - After sudden exposure to pain or an unpleasant sight, sound, emotion, or smell |
| - Long history of recurrent syncope or long duration between episodes (e.g., > 4 years) |
| - Nausea, vomiting, or abdominal pain associated with syncope |
| - Occurs with head rotation or pressure on carotid sinus (e.g., neck tie, tumors, collar) |
| - Prolonged sitting or standing, especially in crowded or hot places |
| - Prandial or post-prandial |

**Syncope due to orthostatic hypotension** |
- After standing up |
- Associated with vasodepressor medications |

**Cardiac syncope** |
- Abnormal ECG |
- During effort or while supine |
- Family history of unexplained sudden death or an inherited condition |
- History of structural heart disease |
- Palpitations followed by syncope |

ECG, electrocardiogram

4.2. Risk stratification schemes

In addition to the above noted clinical findings that help unmask cardiac causes of syncope, and thus identify patients at higher SCD risk, there are several ‘risk stratification’ schemes proposed to aid in clinical decision-making [18,23-31]. Most of these schemes have focused on Emergency Department triage with regard to the need for immediate hospitalization; however, as a rule these schemes only offer short-term risk assessment (typically 1 week to 1 month [Table 3]). The ACC/AHA/HRS 2017 practice guidelines [32] offer a detailed assessment of the available risk stratification tools.

Apart from their short time horizon, caution should be exercised in using these risk stratification tools given the differences in study populations and study designs from which they were derived, and several were not subjected to validation re-testing in a separate patient validation population. Furthermore, it has been suggested that clinical judgment by experienced practitioners performed as well as the prediction tools in an individual patient data meta-analysis identifying serious outcomes at 10 and 30 days after an Emergency Department stay for syncope [33]. In brief, there is no current consensus on which of the risk stratification tools is the most effective; although an ongoing NIH supported study is attempting to address this issue [2].

4.3. Role of ambulatory ECG monitoring

An ambulatory ECG (AECG) is an invaluable tool to diagnose suspected arrhythmias in syncope patients, and the type of AECG chosen depends on the expected frequency of symptomatic episodes [17]. Due to the infrequent nature of symptoms in most patients with recurrent syncope, long-term miniaturized insertable cardiac monitors (ICM) have become widely accepted as a valuable diagnostic tool, especially if an initial wearable AECG monitoring device (e.g., event recorder or Mobile Cardiac Outpatient Telemetry [MCOT]) has been non-diagnostic. The estimated diagnostic yield in syncope is 1–5% for a 24–48 h standard Holter monitor; 5–10% for a 3–7 day patch/external loop recorder/MCOT; 15–25% for a 1–4 week patch/external loop recorder/MCOT; and 30–50% for a < 36 month ICM [34]. Two randomized controlled trials both demonstrated a higher yield with ICM compared to “conventional diagnostic monitoring.” In the first study, ICM was compared with external loop recorder and tilt and electrophysiology testing, with a diagnosis being achieved in 52% versus 20% (at 12 months), respectively [35]. In the second study, the diagnostic yield of ICM versus conventional diagnostics was 43% versus 6% (at seventeen months), respectively [36]. ICM is particularly useful in detecting bradyarrhythmias, which, with rare exceptions (addressed later), are often not detected with an EP study [35]. Importantly, the use of a prolonged ICM monitoring strategy appears to be a safe approach to the diagnostic assessment of recurrent syncope, in most cases.

4.4. EPS in syncope with structural heart disease

An invasive electrophysiology study (EPS) is rarely indicated in syncope evaluation [37]. The greatest diagnostic yield for EPS occurs in those patients with structural heart disease to assess for ventricular arrhythmias, especially in those with prior MI and LV ejection fraction > 35% [37]. Thus, if used in appropriate circumstances, EPS has utility in unmasking those forms of syncope that are at highest SCD risk. Induction of a clinically relevant or hemodynamically significant sustained VT or VF is a class I indication for an ICD [38]. On the other hand, EPS is less useful in the detection of bradyarrhythmias; in certain patients, a His bundle recording can help determine the exact site of a potentially symptomatic AV block. However, long-term ICM is a better choice in this setting, recognizing that some patients may be left at risk of a serious collapse, while awaiting a definitive diagnosis [35].

5. Specific conditions in which syncope has worrisome SCD risk

5.1. Coronary artery disease

CAD remains the leading cause of SCD worldwide [7,39]. Syncope is almost never the cause of myocardial infarction (MI), but is not an infrequent consequence of an acute MI event. One mechanism of syncope associated with myocardial ischemia or acute MI, particularly in the case of an inferior MI, includes a reflex faint with both cardioinhibitory and vasodepressor components presumed to be due to the Bezold-Jarisch mechanism. This latter reflex occurs because of the abundance of mechano- and chemosensitive receptors in the infero-posterior region of the left ventricle, supplied by the inferior coronary vessels, that activate the afferent neural fibers (generally designated C-fibers) of the vagus nerve, and trigger a predominantly vagal-mediated reflex [40]. However, new high-grade AV block on ECG (Fig. 1), or tachyarrhythmias (particularly paroxysmal monomorphic or non-sustained polymorphic VT) are also potential causes of syncope in the setting of acute ischemia. In the primary percutaneous coronary intervention era, among a registry of > 59,000 patients, high-degree AV block occurred in 5.9% of patients with right coronary artery occlusion, and 1.5% of patients with other infarct-related arteries [41].

Particularly concerning is syncope occurring in days to weeks after acute MI. In such cases, EPS may be necessary to unmask susceptibility to life-threatening VT [42]. However, it should be noted that EPS with excessively aggressive stimulation protocols may overestimate this risk due to an increased sensitivity for arrhythmia induction during this period. A typical recommended protocol is ventricular extrastimulus testing at 2 basic pacing frequencies between 600 and 400 ms, and no more than 3 premature
extrastimuli. The additional use of catecholamine infusion can be considered. EPS may be repeated at the right ventricular outflow tract or left ventricle. Limiting the prematurity of the extrastimuli to a minimum of 180 ms is reasonable if sustained monomorphic VT is considered a positive endpoint, as a very short coupling interval is more likely to induce VF compared to monomorphic VT.

In chronic ischemic heart disease, unexplained syncope may be the result of reentrant VT and this circumstance usually merits a

### Table 3

**Principal published syncope risk scores.**

| Study/Author | Sample Size | Outcome Definition | Predictors | Adverse events in lowest risk subgroup |
|--------------|-------------|--------------------|------------|--------------------------------------|
| Osservatorio Epidemiologico sulla Sincope nel Lazio (OESIL) [23] | 270 | 1-year death | Age > 65; CVD in clinical history; No prodrome; Abnormal ECG; | 0% (0% in the validation cohort, n=328) |
| San Francisco Syncope Rule [24,31] | 684 | 7-day serious events | Abnormal ECG; CHF; Shortness of breath; Hematocrit < 30%; SBP < 90 mmHg; | 0.8% (0.3% in the validation cohort, n=371) |
| Boston Syncope Rule [25] | 293 | 30-day serious events | ACS signs/symptoms; signs of conduction disease; worrisome cardiac history; valvular heart disease; family history of sudden death; abnormal vital signs; volume depletion; primary CNS event | 1.4% |
| ECGYS score [18] | 260 | Mortality at mean (SD) follow up of 614 (73) days | Palpitations; Exertional; Supine; Abnormal ECG and/or CVD in clinical history; Autonomic proadrome (negative predictor); Predisposing and/or precipitating factors (negative predictor) | 3% (2% in the validation cohort, n=256) |
| Short-Term Prognosis of Syncope Study (STePS) [26] | 676 | 10-day serious events | Abnormal ECG; Trauma; No prodrome; Male | NA |
| Syncope Risk Score [27] | 2584 | 30-day serious events | Abnormal ECG; Age > 90; Male; Positive troponin; History of arrhythmia; SBP > 160; Near-syncope (negative predictor) | 2.5% |
| Risk Stratification of Syncope in the ED [28] | 550 | 30-day serious events or all cause death | BNP > 300 pg/ml; Fecal occult blood; Hemoglobin ≤ 9; O₂Sat < 94; Abnormal ECG (presence of Q wave) | 0.8% (1.5% in the validation cohort, n=538) |
| Canadian Syncope Risk Score [29] | 4030 | 30-day serious adverse outcomes | Predisposition to VVS; heart disease; any SBP in the ED < 90 or > 180 mmHg; troponin elevation; abnormal QRS axis; QRS > 130 ms; QTc interval > 480 ms; ED diagnosis of cardiac syncope; ED diagnosis of VVS | 0.4% |
| IC-FUC [30] | 393 | 30-day death or unplanned ED/hospital visit | History of heart disease; abnormal ECG; history of syncope | 18.6% |

ACS, acute coronary syndrome; BNP, brain natriuretic peptide; CHF, congestive heart failure; CVD, cardiovascular disease; ECG, electrocardiogram; ED, emergency department; SD, standard deviation; SBP, systolic blood pressure; VVS, vasovagal syncope

*a* Events: death, major therapeutic procedure, myocardial infarction, arrhythmia, pulmonary embolism, stroke, sepsis, hemorrhage, life threatening sequelae of syncope.

*b* Events: death, myocardial infarction, arrhythmia, pulmonary embolism, stroke, hemorrhage, re-admission.

*c* Events: death, major therapeutic procedure, re-admission.

*d* Events: death, arrhythmia, myocardial infarction, new diagnosis of severe structural heart disease, pulmonary embolism, aortic dissection, stroke/TIA, cerebral hemorrhage, significant azemia requiring blood transfusion.

*e* Events: acute myocardial infarction, life-threatening arrhythmia, pacemaker or defibrillator implantation within 1 month of syncope, pulmonary embolus, cerebrovascular accident, intracranial hemorrhage, or subarachnoid hemorrhage, hemorrhage requiring transfusion, or acute surgical procedure or endoscopic intervention.

*f* Events: arrhythmia, myocardial infarction, serious structural heart disease, aortic dissection, pulmonary embolism, severe pulmonary hypertension, severe hemorrhage, subarachnoid hemorrhage, any other serious condition causing syncope and procedural interventions for the treatment of syncope.

**Fig. 1.** 12-lead ECG of a patient with 2:1 AV block. The conducted beats have a long PR interval and a left bundle branch block morphology, indicating severe underlying conduction system disease.
myocardial ischemia evaluation with possible revascularization [37]. Nonetheless, because the substrate for arrhythmia may not be sufficiently modified, despite revascularization, these patients should also be considered for an arrhythmia evaluation, including possible EPS. That said, an ICD may be an appropriate choice, despite a negative EPS in these patients (Class IIb) [38]. Alternatively, an ICM may help in better defining the nature of the unexplained syncope [37]. Both the diagnostic yield and safety of ICM monitoring has been demonstrated in those with syncope associated with structural heart disease [44].

5.2. Idiopathic dilated cardiomyopathy

Syncope is an important risk factor for SCD in advanced heart failure [45], including non-ischemic dilated cardiomyopathy (NIDCM) patients [46]. In such patients, syncope has been shown to have event rates as high as those with sustained arrhythmias [47,48]. Knight et al. followed 14 consecutive patients with NIDCM, unexplained syncope, and a negative EPS. Half the patients had appropriate ICD shocks for ventricular arrhythmias, compared with 8 of 19 (42%) in a control group with prior cardiac arrest (p = 0.1) [47]. Also, Fruhwald et al. followed 23 patients with NIDCM and syncope; 5 of 6 deaths (26% died overall) in the syncope patients were SCDs, whereas only 13 of 41 deaths (20% died overall) in the matched cohort of non-syncope patients were SCDs [46]. For patients with unexplained syncope and NIDCM, it is reasonable to consider ICD therapy according to the most recent guideline recommendations (Class IIa) [38]. Under-recognized causes of NIDCM, such as LMNA-cardiomyopathy, particularly if there are also conduction defects or a family history of NIDCM, must also be considered in those with syncope (Table 4).

### Table 4

Syncope and SCD risk in other cardiac conditions.

| Clinical picture | Refs |
|------------------|------|
| Infiltrative cardiomyopathies (e.g., Amyloidosis, Sarcoïd, Hemochromatosis) | [68,69] |
| Amyloidosis: | |
| – Syncope in ~20% of patients and associated with a poor prognosis | |
| – Multiple mechanisms (postural hypotension most common) | |
| – Cardiac sarcoid and hemochromatosis: | |
| – Conduction abnormalities and ventricular arrhythmias occur frequently | |
| – Unclear if syncope confers a worse prognosis | |
| Early repolarization (ER) | [56,60,70] |
| The finding of ER on ECG after syncope is almost always an incidental finding given the high prevalence of ER in the general population (~5 to 13 percent). Thus, the diagnosis is usually made only after an aborted cardiac arrest or VT/VF in a patient who displays ER in the inferior and/or lateral leads on ECG. | |
| ER syndrome is one of the J wave syndromes and has several similarities with Brugada syndrome. | |
| A case control study of idiopathic VF subjects found that VF cases with ER were more likely to have a history of syncope than VF without ER. Also, syncope may be an important predictor in CPVT patients with ER. | |
| Congenital heart disease | [72] |
| A more aggressive diagnostic approach is recommended when unexplained syncope occurs in congenital heart diseases with “high risk” substrates, including tetralogy of Fallot, TGA after atrial switch surgery, or systemic or single ventricular dysfunction. | |
| Primary pulmonary hypertension | [73,74] |
| There is no consensus on how to interpret syncope in PPH. | |
| Prevalence is more common in children than adults | |
| Multiple mechanisms (e.g., atrial arrhythmias, systemic vasodilation, or extreme transient elevations in pulmonary arterial systolic pressure during exertion) | |
| Laminopathies | [75] |
| LMNA mutation carriers frequently have left ventricular systolic dysfunction and arrhythmias. | |
| In a cohort of 269 Europeans with pathogenic LMNA mutations, unexplained syncope occurred in 11% but was not amongst the independent predictors of malignant VA. | |
| Short QT Syndrome | [55] |
| This rare channelopathy is associated with both syncope and SCD, but whether syncope confers a greater risk of SCD is unclear. | |
| LV Non-compaction | [76] |
| Unexplained syncope has been reported in 5% of LVNC, but it is unknown if this confers an increased risk of SCD. | |

ARIC, atherosclerosis risk in communities; AV, atrioventricular; CI, confidence interval; ECG, electrocardiogram; ER, early repolarization; HR, hazard ratio; LMNA, Lamin A/C; LV, left ventricular; LVEF, left ventricular ejection fraction; LVNC, left ventricular noncompaction; PPH, primary pulmonary hypertension; Refs, references; SCD, sudden cardiac death; TGA, transposition of the great arteries; VA, ventricular arrhythmias; VT, ventricular tachycardia; VF, ventricular fibrillation.
VF and, thereby, be responsible for SCD. Ventricular tachycardia (i.e., torsade de pointe). Torsades may be non-sustained and, in such cases, could be the cause of syncope. However, torsades may also degenerate into polymorphic/bidirectional VT despite seemingly optimal medical management (Class 1) [55]. In CPVT patients with syncope, treatment recommendations start with β-blocker therapy (preferentially nadolol or propranolol) at the highest tolerable dose followed by the addition of flecainide, in the case of recurrent syncope or polymorphic/bidirectional VT while on β-blockers [38,58,59]. Lifestyle changes may also be helpful (i.e., diminishing exposure to excessive exertion). In the setting of failed combination drug therapy, one can then consider ICD implantation, although left cardiac sympathetic denervation has emerged as a promising option [55].

Among 51 patients with CPVT, early repolarization (ER) was present in an unexpected large proportion of 45% (versus the general population of 5 to 13%). A history of syncope was present in 78% of those with ER versus 39% without ER (p=0.005) (Fig. 4) [60].

5.7. Hypertrophic cardiomyopathy (HCM)

Syncope occurs in 15–25% of those with HCM [61]; it is most common in younger patients with smaller ventricles [62], and is often provoked by exercise (during or after) or by postural change. The principal causes of syncope in HCM are broadly divided into arrhythmias and those that result from primary hemodynamic compromise. Fananapazir et al. performed an EPS in 155 patients with HCM, of which 22, 55, and 37 had prior SCD, syncope, and presyncope, respectively [63]. Remarkably, 81% had abnormalities, including sinus node dysfunction (66%), His-Purkinje disease (30%), inducible atrial re-entrant tachycardia (10%), atrial fibrillation (11%), ventricular arrhythmias (43%), and non-sustained VT (14%). Of note, atrial fibrillation is common and may be responsible for clinical deterioration, including syncope and heart failure due to reduced diastolic filling in an already hypertrophic ventricle and reduced cardiac output [61].

Presyncope or syncope in HCM necessitates urgent workup and treatment as such occurrences indicate a high risk of SCD, particularly when recent and occurring in the young. Spirito et al. found that patients with unexplained syncope within 6 months of diagnosis had a 5-fold increase in risk compared to those without syncope [64]. This was most prominent in the young, where those particularly important concern in CPVT in which patients have died despite seemingly appropriate ICD therapy. Thus, ICD is recommended as an early treatment strategy, only in those who have had a definite cardiac arrest, recurrent syncope, or polymorphic/bidirectional VT despite seemingly optimal medical management (Class 1) [55]. In CPVT patients with syncope, treatment recommendations start with β-blocker therapy (preferentially nadolol or propranolol) at the highest tolerable dose followed by the addition of flecainide, in the case of recurrent syncope or polymorphic/bidirectional VT while on β-blockers [38,58,59]. Lifestyle changes may also be helpful (i.e., diminishing exposure to excessive exertion). In the setting of failed combination drug therapy, one can then consider ICD implantation, although left cardiac sympathetic denervation has emerged as a promising option [55].

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< 18 years old had SCD event rates of 293 versus 55 versus 13 per 1000 person-years for recent ( < 6 mo), remote, and no syncope, respectively. In those 18–39 years old, event rates remained high at 46 versus 5 per 1000 person-years for recent versus remote syncope, respectively. In those ≥ 40 years old, event rates were 20 versus 3 per 1000 person-years in those with recent versus remote syncope, respectively.

5.8. Myotonic and muscular dystrophies

The muscular dystrophies are a heterogeneous group of conditions commonly associated with high rates of left ventricular dysfunction and arrhythmias [65]. However, these conditions are often overlooked in the syncope/electrophysiology literature.

Type 1 myotonic dystrophy (DM1) may be the most important of the muscular dystrophies in terms of high rates of arrhythmias and conduction defects, while having a relatively lower incidence of left ventricular dysfunction. In a recent retrospective study of 1388 adults with DM1, sudden cardiac death occurred in 3.6%, accounting for 15.4% of all deaths. Major conduction defects developed in 19.3% and sustained ventricular arrhythmias in 2.3% [66]. Syncope was an independent predictor of major conduction defects and death, but not sudden death or ventricular arrhythmias. Myocardial fibrosis and cardiac-conduction abnormalities are considered to be important mechanisms in the risk of sudden death in DM1, and prophylactic pacing is being used in asymptomatic individuals who have conduction abnormalities on ECG. Abnormal sodium current properties have been implicated in these abnormalities in both human and animal studies, which may, in part, account for the observation that the type 1 Brugada ECG pattern is 50-fold more prevalent in DM1 than the general population. Atrial arrhythmias are also common in DM1, can cause syncope, and are a marker of worse prognosis. In a study of 161 patients with DM1, 27 (17%) had either atrial fibrillation or flutter.

![Fig. 3. 12-lead ECG of a patient with catecholaminergic polymorphic ventricular tachycardia (CPVT). Note that the QRS axis alternates with every other beat, and consequently this arrhythmia is referred to as bidirectional VT. Digitalis toxicity may cause a similar arrhythmia.](image)

![Fig. 4. 12-lead ECG of a patient with early repolarization of the inferolateral leads. This ECG finding has recently been associated with increased SCD propensity.](image)
(AFL), with two presenting with syncope-related AFL with 1:1 AV nodal conduction [67]. These atrial arrhythmias were associated with an increase in mortality (30% died with AF vs 10% without, \( p < 0.01 \)).

5.9. Other causes of increased SCD risk in syncope

A summary of additional cardiac conditions in which syncope may be associated with SCD, but in which the association is less certain or less frequent, is provided in Table 4 [55,56,60,68–76].

6. Syncope in patients with ICD

6.1. Syncope before ICD implantation

A non-randomized registry from the Antiarrhythmics Versus Implantable Defibrillators (AVID) Study assessed patients with a history of unexplained syncope and structural heart disease who were excluded from the main trial due to the absence of VT/VF [77]. Eighty patients in the registry had a positive EPS. This unexplained syncope with positive EPS registry (of which 84% had an ICD) had a similar survival to ICD-treated VT patients from the main trial, both groups being superior to drug-treated VT patients (\( p=0.05 \)). Although the two groups appeared similar in terms of survival, subgroup analyses showed that severe left ventricular dysfunction significantly worsened outcomes in the syncope registry patients, but did not impact the outcomes in ICD-treated VT patients from the main trial. This suggests that syncope itself may not have been the major adverse outcome predictor, but rather LV dysfunction.

In the SCD-HeFT randomized controlled clinical trial, 162 (6%) patients had syncope before randomization [14]. Syncope before randomization was not associated with death (HR 0.98, 95%CI 0.73–1.33; \( p=0.91 \)). However, 38% of patient with syncope before randomization had an appropriate ICD shock compared to 19% of patients without syncope (HR 1.75, 95%CI 1.10 to 2.80, \( p=0.019 \)), suggesting syncope before ICD implantation is associated with an increase in appropriate ICD therapies.

6.2. Syncope after ICD implantation

ICDs, while effective in reducing SCD, may not prevent syncope. Collapse may occur before the device recognizes and terminates the arrhythmia. In a post-hoc analysis from the SCD-HeFT randomized controlled trial, syncope after ICD implantation occurred in 14% of patients, and was associated with an increase in all-cause mortality, cardiovascular mortality and sudden cardiac death despite randomization to an ICD [14]. However, an important limitation of this study was that the causes of syncope in SCD-HeFT were presumptive. Thus, it is uncertain how many patients had arrhythmic syncope. Interestingly, patients with syncope in the SCD-HeFT were more likely to have appropriate ICD shocks, yet ICDs did not protect these patients from dying. One speculation from this post-hoc analysis is the possibility that both syncope and malignant ventricular arrhythmias are markers for a more advanced cardiovascular disease state, and that a life-threatening event in these cases is less amenable to being successfully reversed.

7. Limitations of our understanding

7.1. Temporal relationship between syncope and death – immediate risk or long-term?

Recent syncope is generally assumed to be more concerning with regard to mortality risk than is remote syncope. In this regard, Sheldon et al. suggest that a long history of fainting is more indicative of reflex fains and, consequently, is of lower concern regarding SCD or mortality [78]. However, although the data are sparse, patients with reflex faints may later develop other causes of syncope as they age, and, thereby, become high-risk individuals. Thus, clinicians should not rely on a ‘long history of fainting’ as being indicative of low future risk.

The strongest evidence for recent syncope denoting higher risk is found in LQTS patients. Data from the LQTS Registry found recent syncope to be the most significant predictor of SCD, even greater than the degree of QT prolongation and genotype [79]. In HCM, recent syncope (< 6 mo) is also particularly worrisome, and one could argue that these patients are reasonable candidates for an ICD for primary SCD prevention [64]. On the other hand, recent and remote syncope may not differ in regard to SCD risk in heart failure patients. In a retrospective analysis of 491 consecutive patients with advanced heart failure, no prior cardiac arrest, and a mean left ventricular ejection fraction (LVEF) of 0.20 ± 0.7, of which 12% had a history of syncope, the risk of SCD did not differ between patients with recent (~6 weeks) or non-recent (>6 weeks) syncope (\( p=0.68 \)) [45].

The relationship between syncope timing and mortality risk likely varies among disease states and specific syncope etiology. This relationship warrants further study as its answer may directly impact clinical response to a collapse event.

7.2. Does number of episodes of syncope matter in the risk of SCD?

Multiple syncope events over many years suggests a lower mortality/SCD risk based on the evident survivability of the events. In particular, when structural heart disease has been excluded, patients with multiple episodes of syncope are less likely to have been experiencing life-threatening arrhythmias. Krol et al. evaluated 104 patients with unexplained syncope, of which 31 and 73 subjects had a positive or negative EPS, respectively [80]. Rather surprisingly, a negative EPS was associated with significantly more syncope episodes than a positive study (5.2 versus 2.2, \( p < 0.0001 \)); furthermore, all patients with >6 syncopal episodes had a negative EPS [80]. In another report, \( \geq 4 \) syncopal events in the preceding year was an independent predictor of psychiatric illness [81] suggesting a greater likelihood of pseudosyncope, and a lesser probability of cardiac syncope.

Regarding these studies, conditions in which recurrent syncope may be ominous may have been under-represented due to limitations in sample size. Furthermore, as patients age the risk of more serious forms of syncope increase, and, therefore, someone who may have initially had a benign form of recurrent syncope could coincidentally develop a more serious, unrelated form of syncope.

8. Future research needs

Syncope remains a vexing challenge, despite the clinical research efforts of the past few decades. While many studies have provided important insights into the evaluation and management of syncope, their limitations are manifest in the inconsistency of clinical decision aids and the inability of consensus guidelines to guide the clinician in, not only establishing the cause of the
patient's symptoms, but in also providing an effective treatment strategy and prognosis. The following list offers several topics in which future research may be particularly useful:

a) Syncope in the ED should remain an area of focus given the high cost associated with ED visits, and the impact that ED clinicians have on decisions related to hospital admissions. To better address this problem, Sun et al. have proposed the development of a syncope research agenda that importantly involves multiple specialties and countries [2]. Priorities include 1) appropriately defining syncope, 2) eliminating serious, obvious conditions from future risk prediction tools (e.g., previous studies suggest that a low hemoglobin predicted serious outcomes in syncope, but when patients with obvious bleeding conditions were excluded, hemoglobin level was no longer associated with adverse events), 3) improving diagnostic algorithms for syncope, in part, by adhering to standardized reporting guidelines based on agreed-on data definitions, and 4) determine more specific guidance for the patient's disposition and plan for post-ED care, as most current society guidelines simply recommend admission of "high-risk" patients and discharge of those with "low-risk".

b) It is not definitive whether syncope, per se, is the cause of increased mortality. The underlying disease process may be the primary driver. As Kapoor et al. reported [82], patients identified as having syncope secondary to cardiac disease had a much higher all-cause mortality and SCD risk than patients with syncope, not felt to be due to a cardiac etiology. However, Kapoor et al. went further. They had shown that in syncope patients those with a cardiac etiology did poorly. They then examined this from the opposite direction: do cardiac patients with syncope do worse than cardiac patients without syncope? They found no significant difference in all-cause mortality between patients with cardiac disease and syncope when compared to patients with similar cardiac disease and no syncope [83]. Similarly, in the EGSYS-2 follow-up study, Ungar et al. found that the likelihood of early mortality after syncope was related to the underlying severity of the cardiac disease and not to syncope, per se, or the etiology of the syncope [84]. Findings such as these highlight the importance of treating the underlying cardiac condition in patients with syncope complicating structural heart disease.

c) As noted above, based on SCD-HeFT data, syncope in patients already having an ICD is associated with a worse prognosis despite the presence of the device. Given the increasing prevalence of patients with ICDs, this may be an area where future research could have an important impact. The causes of syncope in this group, if defined and better understood, could lead to more effective treatment strategies. For example, if further studies with better monitoring reveal the cause of syncope to be slow ventricular arrhythmias below the device detection zone, or hemodynamically significant arrhythmias causing collapse prior to the onset of device therapy, such scenarios could potentially be amenable to different treatment algorithms with the ICD.

9. Conclusions

Syncope is a common medical problem with many potential causes. The highest mortality and SCD risk occur when syncope is associated with underlying cardiac disease (including channelpathies), and, in particular, when the underlying cause of syncope is determined to be of a cardiac etiology. In this regard, although syncope may be the first manifestation of an underlying cardiac disease, it may also present later as a heralding sign for SCD. It remains far from clear in many scenarios whether syncope is, itself, the driver of increased mortality in patients with cardiac diseases. Current understanding indicates that reduction of mortality in these patients requires aggressive management of the underlying cardiac condition, in addition to whatever specific therapy may be indicated for prevention of future syncope events. A further understanding of the relation between syncope and SCD risk in various heart diseases may help to identify those individuals who will benefit most from diagnostic procedures and therapeutic interventions.

Conflict of interest

Dr. Benditt is a consultant to Medtronic Inc., and Zoll Corp. He is supported in part by a grant from the Dr. Earl E Bakken Family in support of Heart-Brain research.

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