INTRODUCTION

In 1960 Pick and Langendorf published, "Differentiation of supraventricular and ventricular tachycardia." Sixty years later, differential diagnosis of wide QRS tachycardia (WQRST) on the electrocardiogram remains a challenging exercise. Correct diagnosis is important for prescribing appropriate therapy and determining prognosis. Differential diagnosis of wide QRS tachycardia revolves around differentiation between supraventricular tachycardia with aberrant conduction and ventricular tachycardia. Observations such as clinical history, findings of physical examination during tachycardia, AV dissociation, QRS morphology in lead V1 and lead V6, precordial concordance, RS complexes in precordial leads, contralateral bundle branch block during wide QRS tachycardia, R wave morphologies in lead aVR, and ventricular initial/terminal velocity of conduction ratio can help arrive at the correct diagnosis with reasonable accuracy. The observations described here can help arrive at the correct diagnosis of WQRST with both reasonable accuracy and confidence.

METHODS

Data for this review were identified by searches of MEDLINE, PubMed, and references from relevant articles using the search term “Differential diagnosis of wide QRS complex tachycardia.” Only articles published in English between 1940 and 2021 were included.

REVIEW AND ANALYSIS

Electrical impulses that arise above or within the AV node are labeled supraventricular. As the impulse travels at a speed of three meters per second through the AV node and simultaneously down the bundle branches and His-Purkinje system, it typically produces a narrow QRS complex of 80 milliseconds duration. WQRST is characterized by the QRS duration of more than 120 milliseconds at a heart rate of >100 bpm.

ABERRANT CONDUCTION

A block or delay in conduction of supraventricular impulse in any of the bundle branches or His-Purkinje system will prolong QRS duration and...
alter the morphology of the QRS complex resulting in aberrant conduction. It produces distinctive changes in lead V1 on the electrocardiogram: These electrocardiographic alterations may appear similar irrespective of the cause of slow conduction which may be functional (transient) or structural (permanent) due to scar. The speed of conduction will also be slower if the impulse originates in or traverses the myocardium to reach the His-Purkinje system and will therefore produce a wide QRS complex. A VT originating in the left ventricle will produce RBBB morphology and a VT originating in the right ventricle will correspondingly produce LBBB morphology but under these circumstances, the QRS morphology in V1 may not resemble that of classic right or left bundle branch block.

The refractory periods of the right and left bundle are different. The longer refractory period of the right bundle branch predisposes it to conduction slowing or failure, particularly with sudden acceleration of the heart rate as seen in atrial fibrillation. As a result aberrant conduction, a transient functional block, more often exhibits RBBB morphology in lead V1.

Heart rate exerts an "accordion" effect on the refractory period: as the heart rate slows, the refractory period of the conduction system lengthens and as heart rate speeds up, the refractory period shortens. Therefore the cycle length of the preceding R-R interval determines the refractory period of the following beat. However, following a sudden acceleration in rate, the first few beats may still be attuned to a longer refractory period and conduction may partially or completely fail as evidenced by a wider QRS complex due to bundle branch block. Therefore a long cycle length (R-R interval) followed by a short R-R interval that ends with a wide QRS complex often with RBBB morphology and this may continue for the next several beats until the refractory period shortens or the heart rate decreases. There are three types of aberrant conduction.

4.1 | Ashman's Phenomenon

Ashman's phenomenon refers to aberrant conduction due to critical pre-maturity; identical combinations of long and short cycle lengths may manifest as aberrant conduction (Figure 1A). A long R-R interval is followed by a short R-R interval that ends with a wide QRS complex often with RBBB morphology. On termination of the wide complex rhythm there is no compensatory pause.

4.2 | Acceleration and deceleration dependent aberrancy

Acceleration-dependent aberrancy occurs when the heart rate exceeds the capacity of a diseased conduction system. Increased frequency of impulse transmission results in slowed conduction and widening of the QRS complex; LBBB morphology is more common. It is not dependent on the duration of the preceding cycle length (Figure 1B) often seen in patients with underlying heart disease.

Deceleration-dependent aberrancy is observed as the heart rate slows down, therefore longer cycle lengths are associated with aberrant QRS complexes generally exhibiting LBBB morphology. It should be present in at least two sequential QRS complexes (Figure 1C) and should not be the result of ventricular escape rhythm. Conduction must be from the atrium to the ventricle through the AV node and not through the accessory pathway. It is believed to be due to the slowing of the conduction during phase 4 of the action potential (phase 4 aberrancy).

5 | CAUSES OF WIDE QRS TACHYCARDIA (WQRST)

Causes of WQRST include: (1). Supraventricular tachycardia (SVT) with functional or persistent bundle branch block, (2) Ventricular tachycardia (VT), (3) SVT with preexcitation, or (4) Ventricular paced rhythm.

6 | DIFFERENTIAL DIAGNOSIS OF WIDE QRS TACHYCARDIA

Patient's medical history, physical examination during the arrhythmia, a baseline 12 lead electrocardiogram, and an electrocardiogram recorded during the wide complex tachycardia, are essential for establishing a correct diagnosis.

6.1 | Clinical history

History of prior myocardial infarction (MI) and the onset of WQRST after the diagnosis of myocardial infarction, would suggest a
diagnosis of ventricular tachycardia (sensitivity 97% specificity 100%, PPV 98%). History of congestive heart failure (CHF), angina pectoris also carry a high positive predictive value for VT.

6.2 | Physical examination

Heart rate and hemodynamics such as blood pressure are not helpful in determining the mechanism of tachycardia. A normal blood pressure is not incompatible with the diagnosis of VT and a patient with SVT with aberrancy may present with hypotension. Presence of canon A waves during WQRST suggests the presence of AV dissociation. During AV dissociation atrial systole may occur while the tricuspid valve is closed producing a prominent “a” waves (canon A waves) in jugular venous pulse.

The varying intensity of the first heart sound (S1) suggests the presence of AV dissociation. The position of the AV valve leaflets at the onset of systole and the PR interval determines the intensity of the first heart sound. During VT, because of AV dissociation, there is no constant relation between atrial systole (P waves) and ventricular systole (QRS complex). In the absence of an appropriate PR interval, the first heart sound appears softer than normal and muffled. But during capture beats, because of appropriate timing of atrial and ventricular systole, the first heart sound may be normal or louder. During AV dissociation the P to QRS intervals vary and the intensity of the first heart sound, therefore, varies as well.

6.2.1 | Carotid sinus massage, Valsalva maneuver

By increasing vagal tone, these maneuvers may result in transient AV block thus terminating SVT, but may have no effect on VT.

Adenosine is likely to terminate SVT by interrupting AV nodal conduction. In the presence of atrial tachycardia or atrial flutter with 1:1 AV conduction and aberrant ventricular conduction, adenosine may slow the ventricular rate and reveal atrial P waves or flutter waves long enough to establish a diagnosis. Usually, adenosine has no effect on VT.

7 | ELECTROCARDIOGRAPHIC FEATURES

A QRS duration greater than 140 mseconds favors a diagnosis of VT. Right superior (“northwest”) axis is suggestive of VT. Northwest axis results when ventricular impulses originate in the inferior wall or the apical segment of the ventricle. A change in the frontal plane axis of more than 40 degrees when compared to the QRS axis during sinus rhythm is often observed during VT. However, if the QRS axis and morphology during WQRST are identical to that recorded during sinus rhythm, a diagnosis of VT is unlikely. Presence of electrical alternans does not help in differentiating SVT with aberrant conduction from ventricular tachycardia.

7.1 | AV dissociation

In the absence of retrograde conduction of ventricular impulses to the atria, atrial and ventricular electrical activity is dissociated during VT; there are more QRS complexes than P waves. The presence of AV dissociation strongly favors a diagnosis of VT. The sensitivity of AV dissociation tends to be high (100%) while the specificity is low (20%-50%). It may be difficult to identify P waves in the midst of wide QRS complexes especially when the rate of WQRST is high. Although the presence of AV dissociation is highly suggestive of VT, its absence does not exclude VT. A single dissociated P wave

FIGURE 2 (A) Sinus rhythm, left bundle branch block. (B) VT. The QRS morphology and frontal plane axis are identical to that in sinus rhythm.
at the onset of WQRS could be a reflection of AV dissociation and thus suggestive of VT.²¹

Fusion beats result from simultaneous ventricular depolarization by a supraventricular impulse, usually a sinus, as well as a VT beat. A QRS complex produced by fusion is neither identical to a QRS complex due to VT nor to a QRS complex produced by a sinus impulse alone but appears to be a hybrid between the two forms. The R-R interval between the preceding VT complex and the QRS complex of the fusion beat is often the same as the R-R interval of the VT. The presence of fusion beats is highly suggestive of VT (Figure 3).

Fusion beats should be differentiated from progressive aberrancy that may be observed at the onset of wide complex SVT and may be mistaken for fusion beats. Greenstein, et al, noted that the PR interval preceding a QRS complex that exhibits an intermediate degree of aberrancy is longer than the PR interval of a sinus beat, while the PR interval preceding a QRS complex that results from fusion is shorter than PR intervals of sinus beats.²²

Capture beat occurs when a sinus impulse conducts through the AV node and depolarizes the ventricle producing a narrower QRS complex between wide VT complexes. As a result, the R-R interval between the preceding VT beat and the QRS of the capture beat tends to be shorter than the R-R intervals of the VT (Figure 3). The QRS complex of capture beats may appear similar to the QRS complexes observed during sinus rhythm; capture beats are a hallmark of VT and frequent sinus capture may render an otherwise regular VT irregular.²³

### 7.2 | QRS morphology

WQRS due to VT is far more common in older patients (>35 years of age). Griffith, et al, proposed a simple approach: if a firm diagnosis of SVT with aberrancy cannot be established on the basis of QRS morphology in leads V1 and V6 in this age group then the rhythm should be considered VT by default.²⁴

#### 7.2.1 | Right bundle-branch block morphology

A positive triphasic QRS complex in lead V1 (Figure 4A), particularly if normal axis is present, is likely to represent SVT with aberrant conduction. In one study, 57 of the 69 ECG tracings with confirmed SVT with aberrancy exhibited a triphasic QRS complex in lead V1 whereas only 6 out of 65 ECG tracings of VT had a triphasic QRS complex.²⁵

An R/S ratio >1 in lead V6 suggests a diagnosis of SVT with aberrant conduction.²⁶ Monophasic QRS complexes in V1 and an R/S ratio <1 in V6 suggests a diagnosis of VT.²⁷(Figure 4B).

#### 7.2.2 | Left bundle branch block morphology

The QRS complexes in lead V1 are negative in LBBB morphology. In SVT with LBBB morphology due to aberrant conduction, the QRS complex in lead V1 exhibits a narrow (<30 msec) initial R wave and

**FIGURE 3** 12 lead electrocardiogram of VT showing AV dissociation. The P waves are marked by black dots that march through the QRS complexes. The RR interval from the VT beat to the fusion beat (F) is the same as the R-R interval of the adjacent VT beats. Capture beats (C) are narrow and occur early. Preserved Q waves during the ventricular tachycardia. Deep and wide Q waves are seen in V1 and V2 suggestive of old anterior wall myocardial infarction. There is dominant R in aVR and northwest (right superior quadrant) axis.
a sharp, smooth descent (Figure 5A). Notching in the downslope of the S wave in lead V1 and/or V2 and an interval from the onset of the QRS complex to the nadir of the S wave >60 mseconds suggests a diagnosis of VT (Figure 5B), an observation initially made by Josephson, et al. These findings are more likely to be observed in patients with previous anterior wall infarction.

WQRST with LBBB morphology that exhibits a qR or R/S ratio <1 in lead V6 or right axis deviation suggests a diagnosis of VT. If the WQRST does not conform to any pattern of bundle branch block, it is likely due to VT. If the R wave in lead V1 is wider (>30 mseconds) during tachycardia than during sinus rhythm, the diagnosis is likely to be VT (Figure 6), a pattern more commonly observed in patients with old inferior wall infarction. WQRST in which a predominantly negative QRS complex in lead V1 crosses the baseline several times, resulting in a W pattern, is likely to be VT.

7.3 | Precordial concordance

The presence of predominately positive QRS complexes across leads V1 through V6 is positive concordance (Figure 7A) and the presence of negative QRS complexes across the precordial leads is negative concordance (Figure 7B). During concordance, there is no R/S transition. Precordial concordance during WQRST suggests a diagnosis
of VT (PPV 90%–100%, sensitivity 88%). Although precordial concordance is highly suggestive of VT, its absence does not exclude the diagnosis; less than 20% of WQRST due to VT exhibit concordance. Positive precordial concordance may also be observed during WQRST due to ventricular preexcitation in patients with Wolff-Parkinson-White syndrome.

7.4 | Q waves during WQRST

Q waves suggestive of previous myocardial infarction noted on the patient’s baseline ECG that are also present during WQRST favor a diagnosis of VT. Pathologic Q waves, the result of myocardial scar tissue, can result from infarction, sarcoidosis, or Chagas disease; scar tissue facilitates the occurrence of reentrant VT.

7.5 | Width of the QRS complex compared

A diagnosis of VT is suggested if the width of QRS complexes recorded during WQRST is narrower than the QRS complex recorded while the patient is in sinus rhythm. On the other hand, in the case of WQRST due to SVT the width of the QRS is expected to be the same as during sinus rhythm or wider due to slowing of conduction.

7.6 | Contralateral bundle branch block

If a persistent bundle branch block is present on a baseline ECG, contralateral bundle branch block during a WQRST would suggest a diagnosis of VT. For example, if sinus rhythm baseline EKG shows persistent left bundle branch block and during wide complex tachycardia, the QRS morphology suggests right bundle branch block this could only occur during VT (Figure 8). Supraventricular tachycardia with RBBB morphology aberrancy, in this scenario where there is a pre-existing left bundle branch block during sinus rhythm, physiologically and logically would not be able to sustain AV nodal conduction of supraventricular beats.

7.7 | R wave morphologies in lead aVR

A dominant R wave in lead aVR during WQRST suggests a diagnosis of VT (Figure 3). This likely to be seen in VT that originates in the apex or lower third of the ventricular septum, but only moderate sensitivity and specificity have been reported for this finding as a marker for VT. A qR pattern in aVR with a Q wave >40 mseconds or a negative QRS complex with a notch in the downslope of the initial component also suggests a diagnosis of VT.

Dominant R-wave in aVR during sinus rhythm can be seen in patients with left posterior fascicular block, right ventricular hypertrophy, and prior inferior wall infarction.

7.8 | Ventricular initial/terminal velocity of conduction

The speed of conduction is faster when an electrical impulse travels through the His-Purkinje system and slower if it travels through myocardium or scar tissue. If an impulse is traveling through the AV node and His-Purkinje system, as in the case of SVT, conduction velocity will be rapid and the initial component of the QRS complex will exhibit a sharp inscription. When the impulse exits the conduction system and enters the myocardium, the speed of conduction slows, resulting in slurring of the terminal portion of the QRS complex.

![FIGURE 7](A) Positive concordance. All the QRS complexes from V1 to V6 have positive polarity. (B) Negative concordance. All the QRS complexes from V1 to V6 have negative polarity
Conversely, an impulse that arises from the myocardium or scar tissue, as in VT, will exhibit a slurred initial component as it negotiates its way through the tissue but will achieve faster terminal conduction velocity as it enters the His-Purkinje system. The ventricular initial and terminal conduction velocity ratio ($V_i/V_t$) is the vertical excursion (in mV) recorded during the initial ($V_i$) and terminal ($V_t$) 40 mseconds of the QRS complex. A $V_i/V_t > 1$, is suggestive of SVT. A $V_i/V_t < 1$ is suggestive of VT (Figure 9).

7.9 | Change from wide-complex to narrow complex tachycardia

A change in QRS morphology from wide to narrow QRS complexes favors a diagnosis of SVT with aberrant conduction. If none of these parameters were met, a diagnosis of SVT was established. They added that AV dissociation and the presence of PVC’s during sinus rhythm that exhibit morphology similar to that observed during WQRST increased the probability of a diagnosis of VT.

8 | ALGORITHMS/Criteria

In the past 40 years, several algorithms that approach the differential diagnosis of WQRST in a stepwise fashion have been proposed. Algorithms have used the observations described above in various combinations, focusing on a stepwise analysis of the electrocardiogram. However, if the first step of an algorithm indicates a diagnosis of VT, the observer may not search for other evidence. It is therefore essential that a 12 lead ECG of a WQRST be meticulously analyzed for all the above-described observations, both increasing the confidence in the diagnosis and the interpretive skill of the clinician.

Algorithms do not include the important findings of the history or physical examination. In an attempt to simplify the approach, some algorithms exclude some important observations.

8.1 | Wellens criteria

AV dissociation, QRS width >140 mseconds, and left axis deviation to favor a diagnosis of VT. In addition, using the morphologic appearance of QRS complexes in lead V1 and V6 first described by Marriott and Sandler in 1968, Wellens et al, proposed morphologic appearance of QRS complexes in leads V1 and V6 that could help differentiate SVT with aberrant conduction from VT.

8.2 | Griffith criteria

On the basis of multivariate analysis, Griffith et al, suggested a history of myocardial infarction, QRS morphology in leads aVF and V1, and frontal plane axis >40° when compared with the baseline ECG favored a diagnosis of VT. If none of these parameters were met, a diagnosis of SVT was established. They added that AV dissociation and the presence of PVC’s during sinus rhythm that exhibit morphology similar to that observed during WQRST increased the probability of a diagnosis of VT.

8.3 | Brugada algorithm

Brugada and colleagues proposed that the presence of an RS complex in any precordial lead during WQRST in which the interval from the onset of the QRS complex to the nadir of the S wave is >100 mseconds, the presence of AV dissociation and QRS morphology in leads V1 and
V6 are highly suggestive of VT (sensitivity of 99% and specificity of 96.5%)\(^2\) (Flow chart S1). Other studies were not able to substantiate high levels of sensitivity or specificity using this algorithm.\(^3\)

### 8.4 | Vereckei algorithm

Vereckei and colleagues proposed an algorithm that incorporates AV dissociation, the presence of an initial R wave in lead aVR, assessment of the morphology of wide QRS complex for bundle branch or fascicular block, and the estimation of the initial and terminal ventricular activation velocity ratio which, if <1, suggests a diagnosis of VT.

An updated Vereckei algorithm added three observations pertaining to the QRS morphology of lead aVR: 1. an initial R wave in aVR, 2. the presence of an initial R or Q wave >40 mseconds, and 3. the presence of a notch on the descending limb on a negative QRS complex in aVR. Their current algorithm includes a Vi/Vt, but did not include AV dissociation.\(^4\) (Flow chart S2).

### 8.5 | Lead II R wave peak time

Pava et al, noted that the intrinsicoid deflection time >50 mseconds measured in lead II from the onset to the peak of the QRS complex suggested a diagnosis of VT.\(^5\)

### 8.6 | RS/QRS ratio

RS/QRS ratio, which was defined as the ratio of the interval from the onset of the QRS complex to the nadir of the S wave, divided by the QRS width in lead V6, where a ratio of 0.41 apparently differentiated VT from SVT with aberrant conduction. This algorithm was useful for identifying posterior fascicular VT with RBBB morphology from SVT with aberrancy of RBBB morphology.\(^6\)

### 8.7 | Limb lead algorithm (LLA)

Ventricular tachycardia (VT) was diagnosed in the presence of at least 1 of the following: (1) monophasic R wave in lead aVR, (2) predominantly negative QRS in leads I, II, and III; and (3) opposing QRS complex in the limb leads: concordant monophasic QRS in all three inferior leads and concordant monophasic QRS in 2 or 3 of the remaining limb leads with a polarity opposite to that of the inferior leads. The sensitivity and specificity of this algorithm were comparable to the other algorithms described above. It has not been independently evaluated by other investigators.\(^7\)

Vector orientation of these observations suggests that its accuracy will be high if the VT is arising from the inferior wall, lower septum, or apical segment of the myocardium. The use of single observation is unlikely to result in accurate differentiation of ventricular tachycardia from supraventricular tachycardia with aberrant conduction.

### 8.8 | Scoring system

In this approach presence of initial R wave in V1, initial r 40 ms in V1/V2, notched S in V1, initial R in aVR, lead II R wave peak time ≥50 ms, no RS in V1-V6 each was assigned one point, and atrioventricular dissociation if present was assigned two points. Higher scores were more specific for the diagnosis of VT however were identified only in 33% of all the VTs\(^8\).

Differentiation of WQRSST using mathematical models and digitized electrocardiograms or use of special software to analyze the initial activation of the QRS complex in lead V1 and V2 for differentiating VT from SVT with aberrant conduction\(^9\) did not significantly change the accuracy likely to have limited application in day-to-day clinical practice.\(^10\) A comparison of various criteria and algorithms by independent authors found modest diagnostic accuracy, 70%–75%.\(^11\)

### 9 | PITFALLS AND EXCEPTIONS

The clinical history, findings on physical examination during wide complex tachycardia, and a baseline ECG recorded during sinus rhythm may not be available.

Artifacts can be mistaken for WQRSST, particularly if obtained as a rhythm strip in a telemetry or intensive care unit and may lead to additional but unnecessary tests or treatment. Often the peak of the baseline QRS complex marches through the artifact and can be identified (Figure 11). Some pseudo R-R intervals may not be physiologically tenable and the patient is generally asymptomatic. The cause is commonly loss of electrode/skin contact, patient movement, or movement of the telemetry transmitter module.

In hyperkalemia high levels of extracellular potassium around the myocytes result in the slowing of conduction and prolongation of QRS duration. Atrial standstill may also occur with a concomitant loss of P waves.\(^12\)

Antiarrhythmic drugs, specifically class 1C drugs, prolong the QRS duration and alter the morphology of the QRS complex, making a distinction between VT and SVT with aberrant conduction difficult. Tricyclic antidepressants and antiepileptic drug toxicity can also result in VT.\(^13\)

A QRS duration of less than 140 mseconds can be observed in VT arising from one of the fascicles or in the proximity of His-Purkinje tissue. The relative narrowness of the QRS complex can be mistaken for VT with aberrant conduction. VT due to bundle branch reentry can also present with relatively narrow QRS complexes; the presence of AV dissociation may help in making the correct diagnosis.

Atrioventricular dissociation may occur during junctional tachycardia. Aberrant conduction or pre-existing bundle branch block can cause the rhythm to be mistaken for VT.

QRS complex alternans do not help in differentiating the cause of WQRSST. VT can be irregular during the first 30 seconds.

An ECG recorded during an episode of WQRSST can lead to misdiagnosis if the arm leads have been reversed; VT with aberrancy can
be mistaken for VT and P wave morphology, the hallmark for identifying lead reversal, may not be visible. The frontal plane axis may appear to be in the right superior quadrant, and an R wave may appear in lead aVR as a result of lead reversal, leading to an erroneous diagnosis of VT when the WQRST is really SVT with aberrant conduction.

During VT the initial portion of the QRS complex can be mistaken for atrial activity, leading to an erroneous diagnosis of SVT with aberrancy. When multiple surface leads are recorded, identification of QRS complex onset is enhanced by comparison of multiple leads and what appears to be a P wave can be correctly identified as part of the QRS complex.

A poorly synchronized cardioversion of WQRST due to SVT can induce VT or ventricular fibrillation if the shock is delivered in error on the T wave, falsely reinforcing an initial misdiagnosis of WQRST to be due to VT.

An ECG recorded during sinus rhythm may exhibit an R wave in lead aVR, northwest axis, and/or initial slurring of the QRS complex. An R wave in aVR may be present in patients with right ventricular hypertrophy or extreme right axis deviation. The occurrence of SVT with aberrant conduction in these cases can lead to an erroneous diagnosis of VT.

When a pre-existing intraventricular conduction defect is present during sinus rhythm, the criteria for the differential diagnosis of WQRST are of limited value.52-54

Most of the above observations resulted from meticulous analysis of 12-lead electrocardiograms recorded during sustained, regular WQRST. In some studies, these observations have been correlated with intracardiac electrophysiology studies.55 At times a rhythm strip of non-sustained WQRST, or tracings recorded in telemetry monitoring units are presented for analysis and opinion. Inaccurate lead placement, standardization, recording speed, or lack of other data that can be gleaned from a 12 lead electrocardiogram decreases the accuracy of interpretation.

WQRST recorded in late pregnancy may also pose diagnostic limitations. The orientation of the heart and thus the QRS vector, may change due to the elevation of the diaphragm. Idiopathic VT arising from the outflow tract is more common in this age group. Frontal plane axis deviation and a QRS duration of <140 mseconds are common in outflow tract VT. Conventional criteria tend to be less sensitive when it comes to differentiating SVT with aberrancy from idiopathic VT.56

10 | PRE-EXCITED TACHYCARDIA

In the presence of an accessory pathway capable of antegrade conduction, atrial arrhythmias such as atrial fibrillation, atrial flutter, and atrial tachycardia may conduct through the accessory pathway resulting in a pre-excited WQRST (Figure 10). Unlike the AV node, which has a long refractory period and acts as a gatekeeper that prevents the majority of ectopic atrial impulses from reaching the ventricles, accessory pathways may have very short refractory periods that allow the majority of atrial impulses to reach the ventricles. Antidromic AV reentrant tachycardia will produce a WQRST, but the QRS morphology will depend on the location of the accessory pathway along the AV ring and its ventricular insertion site. The QRS morphology will resemble the morphology of the delta wave observed during sinus rhythm as the ventricle is pre-excited by the accessory pathway bypassing the AV node and His-Purkinje system.37,38

11 | VENTRICULAR PACED RHYTHM

A dual-chamber pacemaker programmed in the DDD mode may track atrial activity during SVT and pace the ventricles at that rate (Figure 12). The pacing spikes may be inconspicuous and therefore may mimic WQRST. A chest X-ray and intracardiac electrograms recorded from the device may assist diagnosis.

12 | CONCLUSION

In an attempt to differentiate WQRST due to aberrant conduction from VT, a detailed clinical history, the findings of physical

**FIGURE 11** Artifacts: Artifacts can be mistaken for nonsustained ventricular tachycardia. In this rhythm strip, sinus rhythm QRS complexes are identified by red dots
examination during wide complex tachycardia, a review of medications, meticulous analysis of the 12-lead ECG both during sinus rhythm and during QRS-T while keeping in mind the pitfalls and exceptions, can help arrive at the correct diagnosis with both reasonable accuracy and confidence.57

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CONFLICT OF INTERESTS

The authors declare no conflict of interest for this article.

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FIGURE 12 Ventricular paced rhythm. A dual chamber pacemaker is programmed in the DDD mode that tracks atrial activity during atrial tachycardia and paces in the ventricle (red dots). When the P wave falls during the refractory period the tracking stops. The next beat (arrow) shows atrial and ventricular paced beats.
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