DIAGNOSTIC CHALLENGES IN CHILD VENTRICULAR TACHYCARDIA

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ABSTRACT
Primary cardiac arrhythmias are much less common in infants and children than in adults, their symptoms can be vague and nonspecific. Although real emergencies because of unstable arrhythmias in children are rare, it is important to identify and properly manage these cases. The study presents the case of a child who was admitted for respiratory symptoms, clinical examination discovering the heart rate of 214 bpm. The child became progressively anxious, showing a respiratory effort. The study of electrocardiogram established the diagnosis of anterior fascicular ventricular tachycardia with a tendency to become permanent. The evolution was favorable after the electrical cardioversion and preventive treatment with Verapamil had been performed.

Keywords: ventricular tachycardia, child, electrical cardioversion

CASE PRESENTATION

Patient B.B., aged 6 years and 5 month is admitted for respiratory symptoms: expiratory productive coughing and dyspnea treated at home for 5 days with Klacid and Eurespal. From previous medical history, we discover multiple infections retain the upper airways that did not require hospitalization.

Clinical examination:
- General affected condition T = +1.23 DS, G = +0.87 DS, PC = +1.5;
- Pale skin, Discreet throat congestion;
- Lungs – rough vesicular murmur, rales, bilateral basal, FR= 32r/min, SaO2(-) = 99%;
- Heart – rhythmic tachycardial heart sounds, FC = 214 b/min;

The remaining devices and systems have indicated normal range. The laboratory data have been within normal range.

Due to increased heart rate, it was decided to perform an electrocardiogram (Fig. 1) which indicated FC: 211/min, regular rhythm, QRS axis of - 90 degrees, the length of QRS complexes is 0.13 sec, with regular succession, monomorphic; atrio-ventricular dissociation and the aspect of right bundle branch block associated with left anterior hemiblock.

Stage diagnosis based on the electrocardiogram was difficult to establish. Therefore, the following have been excluded:
- Paroxysmal supraventricular tachycardia anomalously led: although we have the presence of RS (V1-V6) complex, the duration of RS is over 100 ms in precordial leads, atrio-ventricular dissociation in V1, the initial R wave in aVR > 40 ms support the diagnosis of VT.
Atrial tachycardia with wide QRS complexes
Ventricular fibrillation: disorganized ventricular rhythm
Ventricular flutter: sinusoidal aspect through which ORS seems symmetrical on both sides of the isoelectric line.
Polymorphic VT
Accelerated ideoventricular rhythm: it is a monomorphic ventricular tachycardia with frequency in most cases below 120/min, repetitive by accelerated pathological automatism.

Cardio-thoracic radiograph (Fig. 2) showed hilobasally accentuated pulmonary markings and heart with left arch rectitude (mild cardiomegaly CI: 1.3).

Ecocardiography (Fig. 3) showed a ventricular diastolic dysfunction, tricuspid repression 3rd degree, patent oval foramen, ventricular septum contraction asynchrony, ejection fraction 60%, shortening fraction of 31%.

Structural changes of the heart do not represent a starting point of ventricular tachycardia but rather a consequence of electrophysiological changes.

Anterior fascicular ventricular tachycardia with atrioventricular dissociation was diagnosed, with a tendency to become permanent, right bundle branch block with left anterior hemiblock. The patient received antibiotic treatment, symptomatics, adenosine 2.5 mg initially, then 5 mg, Amiodarone 350 mg x 2/day, tachycardia persists after 24 hours under anti-arrhythmic medication. It was decided that cardioversion should be performed as the functional condition of the child worsens progressively: expiratory dyspnea increased, the extremities being cold and sweaty. General anesthesia with 1 mg iv Midazolamum and Revafl 20 mg intravenously, then 50 J biphasic external electric shock aisle in sinus rhythm 88/min were performed. (Fig. 4)

We continued daily monitoring of ECG; the patient receiving Verapamil 40 mg/day. In evolution, under medical treatment, the child’s general condition remained good, maintaining a heart rate of 100/min.

Ekg at discharge showed a sinus rhythm 100 min, QRS axis: +100 degrees. PQ = 0.14 sec. Right bundle branch block, negative T waves in DII, DIII, AVF and in left precordial leads (suggesting a slight decrease in cardiac output due to ischemia).

At check-up admissions, patient’s condition was good, heart rate of 90 bpm, right bundle branch block persisted.
FIGURE 3. EKG during biphasic electrical cardioversion with 50 j

FIGURE 4. EKG at discharge: FC: 100/min, BRD
DISCUSSION

Ventricular tachycardia is very rare in children, but the impact of this condition on prognosis is extraordinarily high. This is especially important as the higher the heart rate, the more reduced is the ventricular diastole, atrial contribution is missing (atrio-ventricular dissociation), leading to impaired myocardial perfusion and then to sudden death (2,3).

Incessant monomorphic ventricular tachycardia is characterized by the presence of ventricular activity sequences with the same morphology becoming permanent, which is impossible to interrupt under medication (4).

It occurs on a pathological layer (congenital heart disease, ventricular hypertrophy, hypertrophic obstructive cardiomyopathy, arrhythmogenic right ventricular cardiomiopathy, acute myocardial infarction, sarcoidosis, Chagas Disease, Brugada syndrome, long QT syndrome) (11) weakening underventricular fibrillation (5) or in a normal heart being very well tolerated, but it may cause in time dilated cardiomyopathy (6).

By Verekei’s algorithm, we can differentiate a ventricular tachycardia from a paroxysmal supraventricular tachycardia by:

• the presence of an initial R in aVR; where initial r or q duration is over 40 ms in aVR; “notched” view of initial slope of the QRS complex predominantly negative in aVR
• ventricular activation-velocity report (Vi/Vt) ≤ 1 (slope of first 40 ms of the QRS complex of aVR/slope of the last 40 ms of the QRS complex of aVR) (7).

If the patient’s condition is stable and is not accompanied by hemodynamic deterioration after proper diagnosis of ventricular tachycardia, cardioversion is attempted with Amiodarone 5 mg/kg intravenously or Procainamide 15 mg/kg intravenously (Amiodarone and Procainamide are never administered together) or Lidocaine 1 mg/kg bolus.

If drug cardioversion was not successful, then the synchronous biphasic electrical cardioversion synchronous is used of 0.5-1 J/kg (2 J/kg may be attempted if the initial dose was ineffective); these maneuvers are done only after prior sedation.

Postcardioversion, in fascicular ventricular tachycardia, relapse prevention is achieved by administering orally Verapamil (9,10). If symptoms reappear, or there is a drug-intolerance, invasive treatment methods may be applied:

• Implantable cardio-defibrillator (if there is an increased family risk of sudden death: Long QT syndrome, Brugada syndrome, ventricular tachycardia with severe hemodynamic and ejection fraction <35%);
• Radiofrequency ablation treatment (especially indicated in normal cardiac ventricular tachycardia in certain diseases but also in specific pathologies: fascicular tachycardia, idiopathic tachycardia, arrhythmogenic right ventricular dysplasia) (8);
• ablative treatment by other forms of energy: cryoablation, microwaves, laser and ultrasound.

CONCLUSIONS

Ventricular tachycardia may be an individual pathology on normal heart differently constituted organically or structurally. Good tolerance of symptoms and lack of proper cardiac abnormalities that could have led triggered ventricular tachycardia led to the establishment of anti-arrhythmic medication but were not therapeutically successful. Electrical cardioversion determine the restoring of electrographic path and the normalization of heart rate with a long-term favorable outcome.
REFERENCES