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# BLOQUEO AURICULO VENTRICULAR.

## 1. Sistema de Conducción Cardíaco

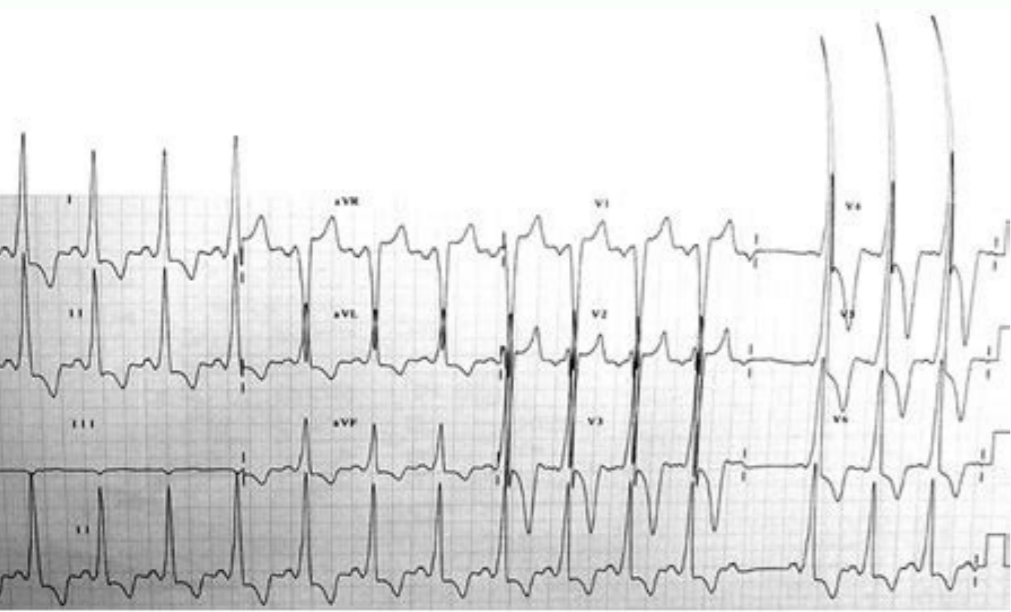


Figura 1. Electrocardiograma de superficie que muestra intervalo de PR corto, complejo QRS prolongado y onda delta.

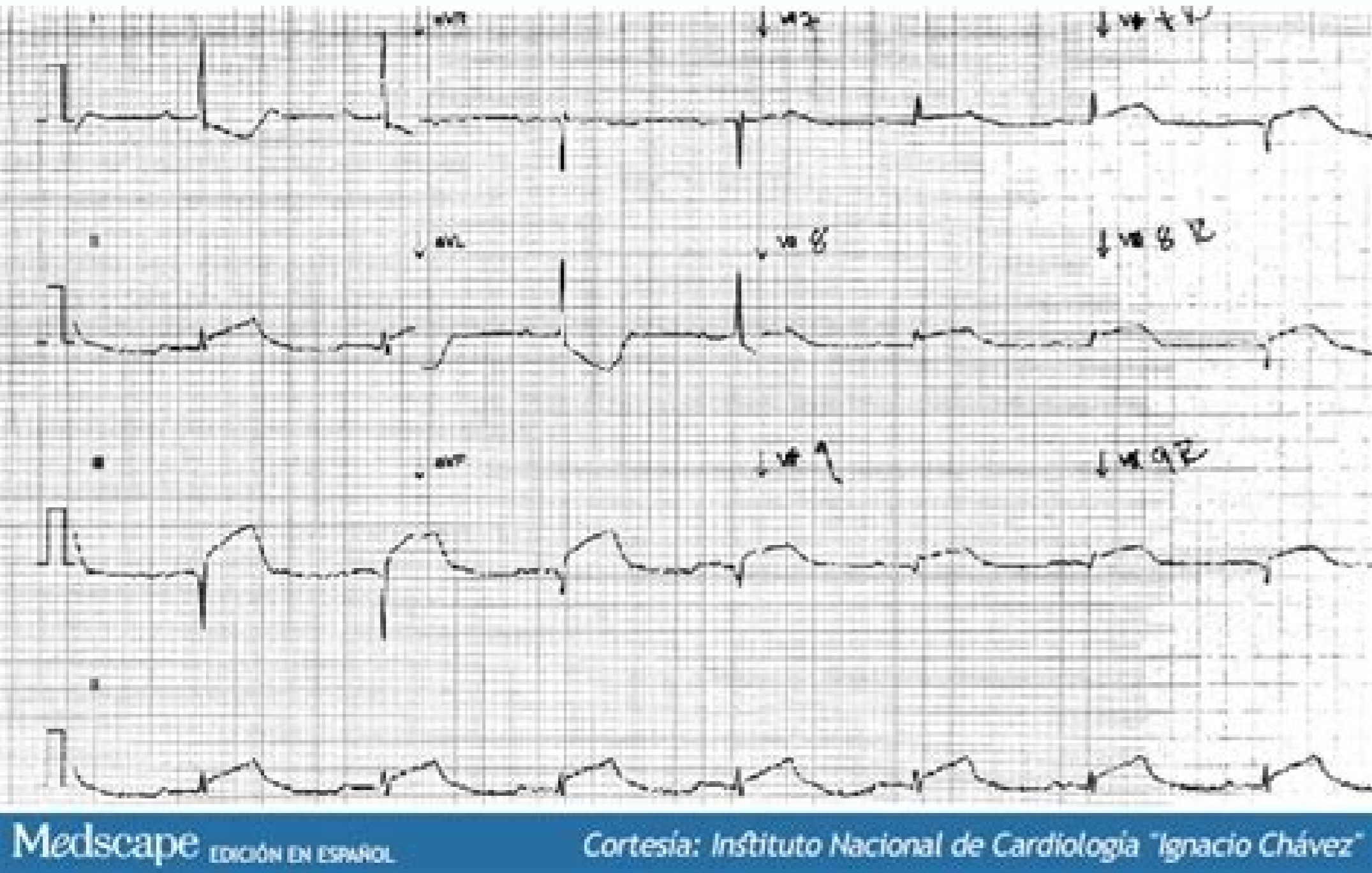
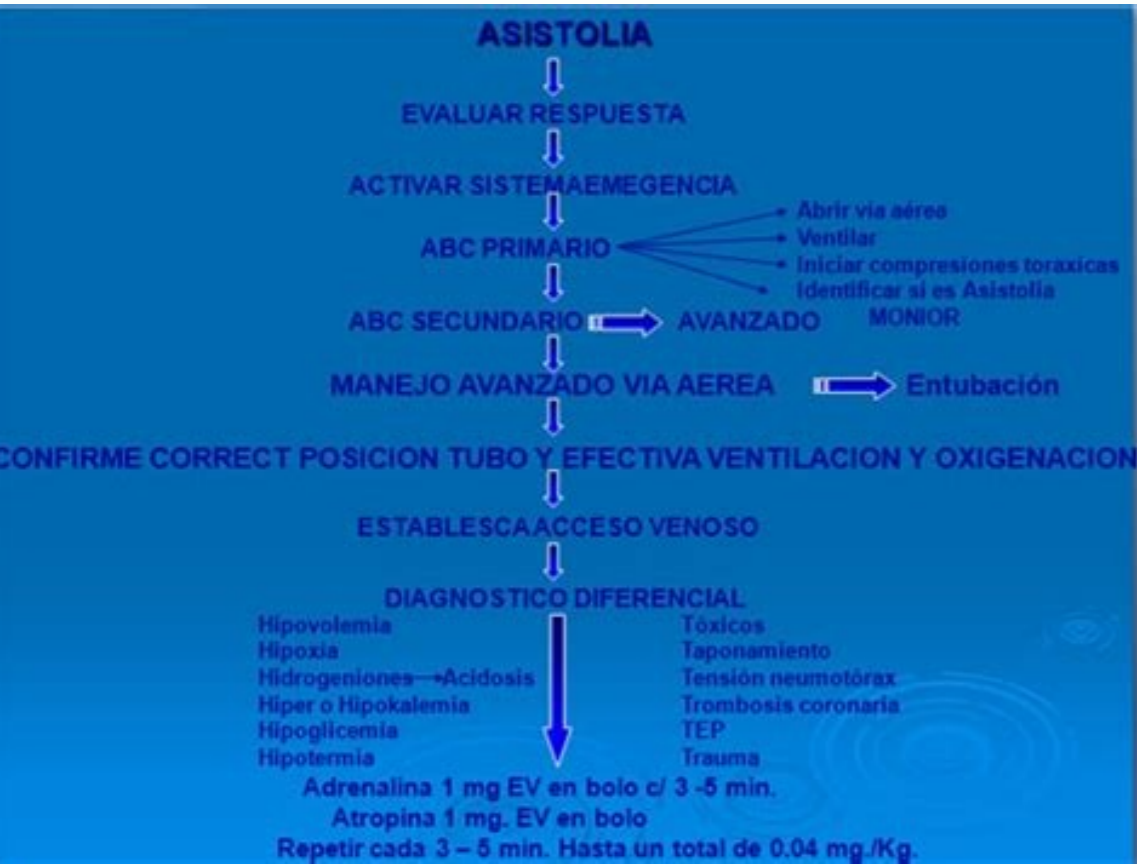


Table with 2 columns: 'Bloqueo Auriculo Ventricular Clasificación' and 'Clasificación'.

Recently, Zizek et al.14 implemented a resynchronizer in a patient with primary amyloidosis and administered chemotherapy, finding better in the function and ventricular volumes at the 12 months of follow-up. CONCLUSIONS A patient with cardiac amyloidosis is initially manifested as paroxysmal AV block with the commitment of left ventricular function and heart failure in functional class. The one of these patients is a challenge for closing management, given the little tolerance to several cardiovascular medications, and the cornerstone of treatment is the recognition of its primary or secondary etiology for adequate management according to its cause. Rudolf Virchow describes the "amyloid" (from the Greek Amylon, "starch"), observing its similarity with the iodine and Edo sulfá, sulférico1. This suggests that cardiac commitment at the conduct system level is very frequent, as manifested in our patient, and should be routinely evaluated in any patient with amyloidosis, through the taking of an electrocardiogram in the asymptomatic patient or of more complete forms in cardiovascular symptoms (Holter, event monitor or electrophysiological study). When there are technical limitations in the imagiography blood and myocardium9. Confirmation of the diagnostic cardiac amyloidosis requires the demonstration of amyloid depth in a biopsy, although it does not necessarily have to be cardiac. When withdrawal of the device is found with sinus bradycardia without advanced AV block (Fig. Our case did not present positivity of any of them. Múltiples have been tested Modifiers of evolution of the disease (D-Penicilamine, immunosuppressants, cyclosporine, etc.) and none to date has been effective in double blind tests3. Among other anomalies, ventricular arrhythmias, which are frequent, although they rarely produce syncope and do not usually be the symptom of presentation symptoms. The abnormalities in the conduct system and arrhythmias are frequent2, two). However, there is reported that patients with amyloidosis had marked dissynchronia, although it has not been reported by other authors. This is a dominant AutoMie disease with a high risk of death of which even the prophylactic use of pacemaker is raised. The beats are not conducted intermittently and the complexes disappear, in general in repeated cycles every three (blocking of 3: 1) or four (block of 4: 1) P waves. The risk of complete av block is differ Predict and is usually indicated. It is also observed fibrilación atrial due to the progressive dilatation of the auritas and the restrictive physiology. The presence of antibodies in these patients has been studied and it has been reported that 33% have antinucleares Those presented antipolysse I, 75% have an ECS and 52% affectionate cardíaca5,6. Radiography and axial computed tomography of Tájá\*rax targeted a lung -nquima with signs of chronic obstructive pulmonary disease and past tuberculous disease. Echocardiografía\*ba objectivizer left ventricular hypertrophy with disfunciá\*ba n diast. The sclerosis of the peribulbar grooves can be seen. The complementary tests showed as relevant data: creatinphosphocinase of 1,545 IU/L with fraction MB of 80, lactate dehydrogenase of 865 IU/L, transaminase glutá\*amico oxalacó 58 IU/L, glutá\*amico pirá\*ávica de 63 UI/L, gammaglutamiltranspeptidasa de 46 UI/L, aldolasa 12 UI/L (normal de 0,3-7,6); leucocitos de 13.600/mm3 (con 15% de linfocitos y 69% de neutró\*filos); y velocidad de sedimentaci\*An globular de 14 mm. Electrocardiograma que demuestra el bloqueo auriculoventricular completo.Si bien la afectaci\*An cardíaca por la ECD es bien conocida y se presenta con una prevalencia del 1-2%, su debut como BAVC es rara. Al examen fásico presenta hipotensi\*An (80/40), FC 64/min, ingurgitaci\*An yugular GII, disimuci\*An bibasal del murmullo vesicular sin estertores, con ascitis y edema gradoIII en miembros inferiores. En estudios adicionales se descarta amiloidosis primaria con biopsia de la má\*Adula a\*Ásea, la electrofresis de proteí\*Anas, la serie a\*Ásea y la biopsia de grasa subcutá\*Ánea. En el seguimiento a los 3 meses, el paciente se encuentra en clase funcionalIII-IV, no ha presentado pres\*Ancope o s\*Ancope. Los betabloqueadores son a menudo evitados por la misma raz\*An. Es una enfermedad poco conocida y muchas veces subdiagnosticada, en la cual aun contando con el diagn\*Ástico de compromiso cardíaco, es necesario aclarar su origen debido a las má\*Átiples causas e implicaciones en su manejo y pron\*Ástico.Caso clá\*ÁnicoPaciente de 69a±Aos de edad que asiste al servicio de urgencias por deterioro de la clase funcional, con disnea clase funcionalIV, ortopnea, disnea paroxá\*Ástica nocturna y edema de miembros inferiores. Se da tratamiento antibi\*Ático con ceftriaxona. Tiene una afectaci\*An má\*Ás limitada, infiltrando el ri\*Á\*An, las suprarrenales, el h\*Ágado y el bazo5.La amiloidosis hereditaria o relacionada con la transtirretina se debe a mutaciones de la Transtirret(An. 4), no presenta episodios de bajo gasto o s\*Ancope. Ingres\*Á en la Unidad Coronaria, donde se le coloc\*Á un marcapasos definitivo tipo VVI-R que mejor\*Á la clá\*Ánica que motiv\*Á a su consulta. Es una causa importante de las enfermedades They can be responsible for both cardiac commitment and other O\*rganos, generating malfunction of these2,3. They have been identified in the histology characteristic as their polychromatism with violet crystal, the violation of Schiff's periodic periodic, although the most used histological criterion is used is the positivity for the Congo Red (observing a greenfringence Apple characteristic in the study of polarized light) 1-3. Amiloidosis can be classified according to 2 criteria: á € according to the distribution of amyloid depths, in localized or systemic forms. The 4 main types of systemic amyloidosis are: primary amyloidosis, family amyloidosis (or hereditary amyloidosis), secondary amyloidosis (reactive amyloidosis) and systemic amyloidosis seni4. Primary or light chain amyloidosis has an incidence of approximately one case for every 100,000 people in the western countries. It was performed a physical exploration that highlighted: atóric, resisted, tense, tense skin, which affected a trunk (toric sclerosis), face and limbs, thickening of the peribulbar grooves (Fig. It is associated with the disagreement of the C\*Á Lulas plasma, being the main mylom Cresil). Myocardial fibrosis is the open finding of the ECD with cardiac involvement, detected by gammagraph of perfusion with talium1. Cardiac affection is found in 9.9% of patients after 5.7 years of follow-up follow-up with a 12% mortality for this cause4. A study carried out in 163 patients with cardiac commitment for amyloidosis found the presence of auriculares en el 15.9%, bajo voltaje en derivaciones de las extremidades en el 54.5% e imagen de pseudoinfarto en el 40.2%8. Habitualmente son sintomá\*Áticos y de evoluci\*An fatal sin tratamiento. El electrocardiograma que se le realiz\*Á al ingreso presentaba un BAVC, con complejo QRS ancho a 35 latidos por minuto (fig. Si la biopsia de otros tejidos es negativa y la sospecha persiste, la biopsia cardíaca es necesaria8,9. Describimos un caso de un paciente de 69a±Aos de edad, con amiloidosis que ingresa al servicio de urgencias con un bloqueo auriculoventricular completo.Cardiac amyloidosis is a manifestation of a systemic diseases group characterized by protein misfolding caused by extracellular deposition of amyloid, known as amyloidosis. 1). Introducci\*AnLa amiloidosis cardíaca es una forma de manifestaci\*An de una serie de patologí\*As sist\*Ámicas. Copyright Á© 2015. Cortesí\*An de Algunos ondas P normales presentan complejos QRS a continuaci\*An de ellas, pero algunas no lo hacen. Ha sido revisado periá\*Ádicamente y tras 2 a\*Áos de seguimiento presenta mejorí\*An clá\*Ánica moderada. Nuestro paciente se present\*Á como falla cardíaca asociado a bloqueo AV.En el ecocardiograma se ha descrito el engrosamiento de las paredes de los ventr\*Áculos de predominio izquierdo, la disfunci\*An diastá\*Álica, la dilataci\*An auricular, la presencia de un patr\*Án a\*Ágranular a del miocardio y el derrame pericá\*Árdico. Sociedad Colombiana de Cardiología y Cirugía Cardiovascular Se le realiza un estudio de biopsia endomiocá\*Árdica documentando en la histología\*Ámica como reactividad en el material amorfó extracelular para el rojo Congo alcalino, confirmando que se trata de la amiloide (figs. Los marcapasos pueden ser necesarios en funci\*An del grado de la enfermedad del sistema de conducci\*An10. Al ingreso se encuentra al paciente con PA 90/40, FC 72/min, FR 16/min. Un comentario aparte merece la amiloidosis familiar con polineuropatía, la it is associated very commonly to conduct disorders and sinus dysfunction13. Cardiac commitment in these patients is pronostico and can be manifested as severe deterioration of the FEVI associated with heart failure or, as in our patient, by commitment of conduction av. Visceral and cardage affection in particular is more frequent in patients with ECS. Asymptomatic arrhythmias are very common in patients with ECS. The presence of heart disease overwhelms the pronostico of this entity. We contribute the case of an ECD associated with a third-grade or complete aurricular blockade (BAVC) that, although it has been documented in the literature, is rare. He did not have telangiectasias, neither digital á \*ñceras, nor crashed pulpejos scars, nor apparent calcinosis. The patient has a previous background a month ago of cardioresinronizer cardiore in another institution by full AV block (Fig. In our patient, ECA or beta blockers could not be used by hypotension and a history of bradyarrhythmia. The pronostico for patients with primary amyloidosis is poor . Cutaneous biopsy was practiced and a marked marked fibrosis was appreciated with actotic elastosis and absence of hairicious folacles compatible with sclerodermia. They are clogically asymptomatic in its majority. Cardiac amyloidosis is a manifestation of a group of disease micas that is characterized by the bad folding of protein that causes an extracellular deposit of the amyloid, known as amyloidosis. In our patient there was improvement of the fraction of ejet Randomized double blind study has been unable to demonstrate any benefit in The recent ECS beginning7. 2 and 3). He took coronary arteriography with evidence of healthy epic arteries. The It is a group of systemic diseases that is characterized by the bad folding of the proteins that causes extracellular situation of fibrillar protein material, insoluble and resistant to the action of proteolytic enzymes.



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