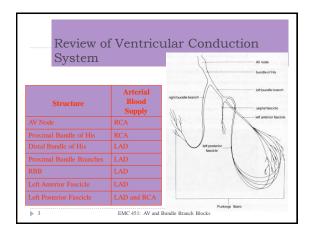
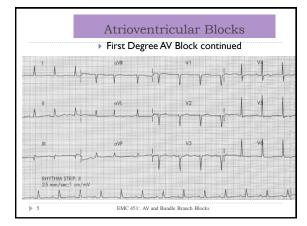


# Unit Objectives Upon completion of this unit, the student will be able to: Describe the normal ventricular conduction system. Recognize the ECG changes associated with 1, 2, and 3 AV block. Discuss the clinical significance of 1, 2, and 3 AV block. List the instances of AV block when pacemaker insertion is necessary. List the instances of AV block when the use of atropine is contraindicated. Recognize the ECG changes of RBBB. Discuss the clinical significance of RBBB. Recognize the ECG changes of LBBB. Discuss the clinical significance of LBBB.



# Atrioventricular Blocks First Degree AV Block Not really a block Delay in conduction through either the AV node or His bundle. PR interval > 0.20 seconds PR interval usually < 0.40 but may be up to .60 seconds Rarely, PR intervals as long as 1.0 seconds (5 large boxes) have been recorded



### AV Blocks continued

### First Degree AV Block continued

- ▶ Clinical Significance
  - Common finding in normal hearts (3% incidence)
  - Occurs in up to 13% of patients with AMI
  - May be an early sign of degenerative disease of the conduction system
  - $\,\,\,\,\,\,\,\,\,$  May be transient sign of myocarditis or drug toxicity
  - Clinical significance is not a function of the amount of PR interval prolongation
  - Rarely requires treatment.

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### AV Blocks continued Second Degree AV Block Type I (Wenkebach) Almost always due to a block within the AV node Increasing PR interval until a beat is dropped 2° AV Blocks Sequence repeats itself, but the Wenckebach ratio of conducted to nonconducted beats does not have to remain constant ▶ The block may be intermittent R-R interval progressively His Bundle shortens within each group of Right Bundle Branch conducted beats Left Bundle Br EMC 451: AV and Bundle Branch Blocks

### AV Blocks continued

Second Degree AV Block Type I (Wenkebach) continued

- QRS complexes are narrow unless there is an associated BBB
- Clinical Significance
- Usually does not progress to more advanced conduction problem
- Commonly associated with digitalis toxicity, inferior wall MI, RV MI, and acute myocarditis
- May also be a normal finding in trained athletes
- If complete heart block develops because of inferior wall MI, the escape mechanism is junctional and does not always necessitate pacemaker insertion.

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EMC 451: AV and Bundle Branch Blocks

# AV Blocks continued Second Degree AV Block Type I (Wenkebach) continued

### Second Degree AV Block Type II Usually due to a block below the AV node, within or below the His bundle Almost always associated with RBBB No lengthening of PR interval, or shortening of RR interval, as with Wenkebach 2° AV Blocks ▶ P waves conducted at 2:1-4:1 ratios Wenckebach commonly, but higher ratios are possible P to R conduction ratios may vary QRS complexes usually broad. The complete block of one bundle causes His Bundle the ORS to be broad, while the intermittent block of the other bundle causes the dropped beats. EMC 451: AV and Bundle Branch Blocks 10

### AV Blocks continued

### Clinical Significance

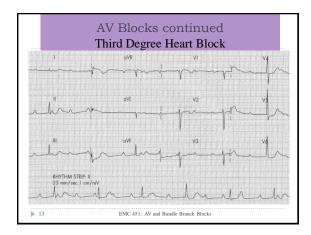
- Not as common as Type I AV block, and is more serious.
- Denotes disease in lower portions of the conduction system.
- Often associated with Stokes-Adams syncope and deteriorates into complete AV block.
- Associated with anteroseptal MI and chronic fibrotic disease of the conduction system.
- Indicates a high risk patient when associated with anteroseptal MI.
- When complete heart block develops, the escape mechanism is slow and of ventricular origin.
- Atropine is rarely effective in improving ventricular rate, thus requireing pacemaker insertion.
- Impossible to differentiate Type II with 2:1 conduction

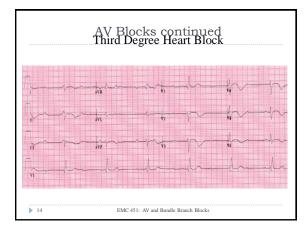
from Wenkebach with 12 lead EKG.

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### AV Blocks continued Third Degree Heart Block Forms of Complete (3°) AV Block No atrial impulses are conducted Block may be located in the AV node, His bundle, or bilateral BBB Complete block in the upper AV Node leaves Junctional foci to Complete block of the entire AV Node or in the His Bundle leaves Bundle, all paths If block in upper AV node, Node leaves Junctional foci to escape and pace the ventricles. are completely blocked, so a ventricular focus junctional escape rhythm ensues, otherwise, escape rhythm is only a ventricula focus to pace. ventricular and ORS is wide. 12 EMC 451: AV and Bundle Branch Blocks

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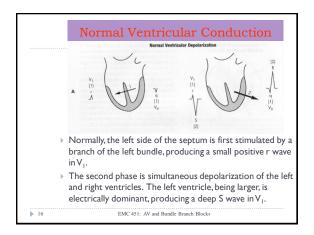


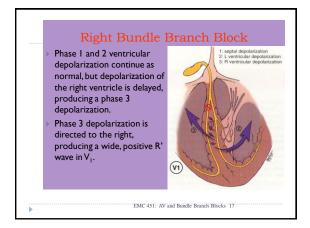
### **AV Blocks continued**

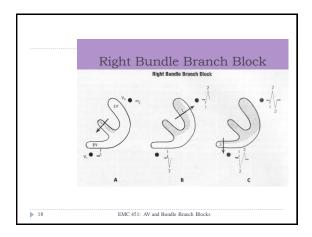
### Clinical Significance of 3<sup>rd</sup> Degree Heart Block

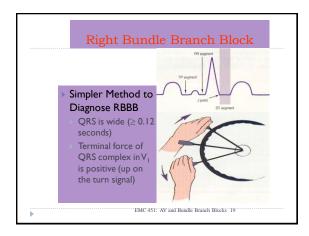
- Rate and dependability of the escape rhythm is related to the level of the lesion.
- An escape at the top of the His bundle has a rate of about 55 bpm, is relatively dependable, and may not require pacemaker insertion. This type of 3<sup>rd</sup> degree block is associated with inferior MI.
- Lower level lesions produce wide QRS complexes, slower rates, less dependable escape rhythms, and require pacemaker insertion. These are usually associated with anterior MI.
- ANY dissociation loses "atrial kick," reducing cardiac output that may result in syncope or angina. If the block is below the AV node, increased SA firing with atropine, exercise, or catecholamines would compound the block. If the block is at the AV node, increased SA firing and AV conduction speed would improve the block.

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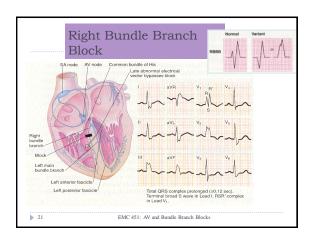


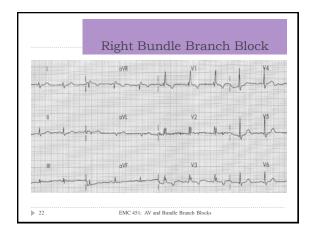
### Right Bundle Branch Block and T Wave Changes

- Because ventricular depolarization is abnormal, so are the T waves. The T wave is expected to be opposite in polarity to the terminal component of the QRS complex in the right precordial leads. This is known as secondary T wave changes and is a normal consequence of BBB.

  If the polarity of the T wave is the same as that of the terminal component of the QRS complex, myocardial disease is suspected.
- T wave changes in leads other than V, are primary T wave changes and are not the result of RBBB.

EMC 451: AV and Bundle Branch Blocks





### Right Bundle Branch Block Clinical Significance May be caused by a number of factors, including COPD, atrial septal defect, pulmonary hypertension, pulmonary stenosis, pulmonary embolism, anterior wall MI, inferior wall MI, and degenerative changes of the conduction system. When associated with anterior wall infarction, new onset RBBB indicates a risk for complete heart block, especially when accompanied by left anterior or posterior hemiblock. When associated with anterior wall MI, mortality is high because it signifies an orclusion in the proximal LAD.

When associated with anterior wall MI, mortality is high because it signifies an occlusion in the proximal LAD. Even without complete heart block, mortality is high because of extensive myocardial damage and pump failure. Normally, RBB does not require treatment.

EMC 451: AV and Bundle Branch Blocks

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Left Bundle Branch Block

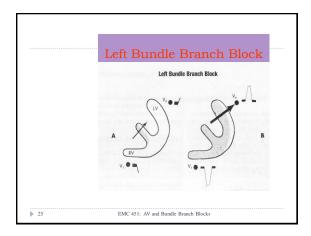
Normally, the septum depolarizes from left to right. But in the presence of LBBB, the septum depolarizes from right to left.

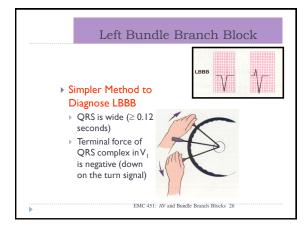
Results from either total occlusion of left main branch or a simultaneous block in both anterior and posterior fasicles.

The normal septal r wave in V<sub>1</sub> is lost, and the total time for left ventricular depolarization is prolonged.

A QS pattern is produced in V<sub>1</sub>.

Axis is normal or deviated to the left.

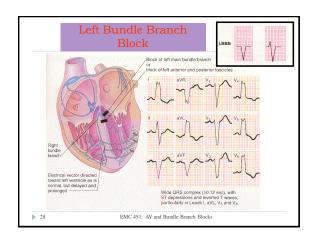


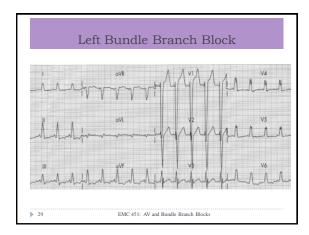


## Left Bundle Branch Block • Because ventricular depolarization is abnormal, so are the T waves. The T wave is expected to be opposite in polarity to the terminal component of the QRS complex in the left precordial leads(V<sub>5</sub> – V<sub>6</sub>). This is known as secondary T wave changes and is a normal consequence of BBB. • If the polarity of the T wave is the same as that of the terminal component of the QRS complex, myocardial disease is suspected. • T wave changes in leads other than V<sub>5</sub> – V<sub>6</sub> are primary T wave changes and are not the result of LBBB

EMC 451: AV and Bundle Branch Blocks

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# Usually a sign of organic heart disease (ischemic LCA disease, chronic hypertensive heart disease, calcification of the mitral valve, aortic stenosis or regurgitation, and cardiomyopathy) Most patients with LBBB have underlying LVH and impaired LV function. Degenerative changes in the conduction system may lead to LBBB, particularly in the elderly. LBBB may be the first clue to 4 previously undiagnosed pathologies: CAD, valvular heart disease, hypertensive heart disease, and cardiomyopathy. It is very difficult to diagnose AMI in the setting of LBBB. MEMC 451: AV and Bundle Branch Blocks