

## ANTIARRHYTHMIC PHARMACOLOGY QUIZ%0A

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Antiarrhythmic agents differ in their effects based on their pharmacologic characteristics. Some medications influence ion channels in conducting cells (ex: SA and AV node), while others influence ion channels in working myocardial muscle cells.

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### **Medical Pharmacology Chapter 8 Antiarrhythmic Drugs**

A patient is diagnosed with atrial fibrillation and exhibits a ventricular following rate of 150 beats / min. Quinidine gluconate (Quinaglute, Quinalan) was administered and although atrial fibrillation was converted to atrial flutter, the ventricular rate increased to 190 beats / min.

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### **Antiarrhythmic Drugs PharmaFactz**

In this guide, we examine the mechanistic pharmacology of antiarrhythmic drugs. Later articles detail clinical aspects side effects and drug interactions of each class. It s important that the student understands how these drugs work on a mechanistic level, before immersing in the many clinical factors behind each medicine.

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### **CV Pharmacology Antiarrhythmic Drugs**

All antiarrhythmic drugs directly or indirectly alter membrane ion conductances, which in turn alters the physical characteristics of cardiac action potentials. For example, some drugs are used to block fast sodium channels. These channels determine how fast the membrane depolarizes during an action potential.

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### **Chapter 45 Antiarrhythmic Agents My Nursing Test Banks**

Quinidine is a class I antiarrhythmic and stabilizes the cell membrane by binding to sodium channels, depressing phase 0 of the action potential, and changing the duration of the action potential. Disopyramide, moricizine, and procainamide are all used in the treatment of life-threatening ventricular arrhythmias.

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