

Federal Employee Program.

TIKOSYN (dofetilide)

RATIONALE FOR INCLUSION IN PA PROGRAM

Background

Tikosyn (dofetilide) is an antiarrhythmic drug with Class III (cardiac action potential duration prolonging) properties. Tikosyn blocks cardiac ion channels carrying the rapid component of the delayed rectifier potassium current I_{KR} . At clinically relevant concentrations, Tikosyn has no effect on sodium channels, adrenergic alpha-receptors, or adrenergic beta-receptors (1).

Regulatory Status

FDA-approved indication: Tikosyn is indicated in patients with atrial fibrillation/atrial flutter (1).

Summary

Tikosyn (dofetilide) is and antiarrhythmic drug with Class III (cardiac action potential duration prolonging) properties. Tikosyn blocks cardiac ion channels carrying the rapid component of the delayed rectifier potassium current I_{KR} . At clinically relevant concentrations, Tikosyn has no effect on sodium channels, adrenergic alpha-receptors, or adrenergic beta-receptors (1).

Prior approval is required to ensure the safe, clinically appropriate and cost-effective use of Tikosyn while maintaining optimal therapeutic outcomes.

References

1. Tikosyn [package insert]. New York, NY: Pfizer Inc.; July 2021.