Metoprolol (Lopressor, Toprol-XL) Considerations for Use*

US/FDA Approved Indication: Heart Rate Control for Atrial Fibrillation

| Black Box Warning* | Abrupt cessation may exacerbate angina pectoris and MI. |
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| Mechanism of Action | Blocks binding of catecholamines to beta-1 receptors; Beta-1 selective |
| Dosing [†] | Acute setting: 2.5 to 5 mg IV bolus over 2 minutes; may repeat every 5 minutes to a maximum dose of 15 mg |
| | Non-acute setting or maintenance: Metoprolol tartrate (immediate-release): 25 to 100 mg PO twice daily |
| | Metoprolol succinate (extended-release): 25 to 200 mg daily |
| | Elderly: May need lower doses |
| | Hepatic Impairment: May need lower doses |
| | Renal Impairment: No dosage adjustment needed |
| Contraindications | AV block |
| | Bradycardia |
| | cardiogenic shock |
| | decompensated heart failure |
| | sick sinus syndrome |
| | pheochromocytoma |
| Major Side Effects | hypotension, heart block, bradycardia, bronchospasm, HF |
| Dosage forms and Strengths | <u>PO:</u> |
| | 25 mg, 50 mg, 100 mg immediate-release tablets (tartrate) |
| | 25 mg, 50 mg, 100 mg, 200 mg extended-release tablets (succinate) |
| | <u>IV:</u> 5 mL ampules (1 mg/mL) |
| Special Notes | Abrupt cessation my precipitate angina, MI, arrhythmias, or rebound HTN; discontinue by tapering over 1-2 weeks. |
| | Immediate-release form is metoprolol tartrate; extended-release form is metoprolol succinate. When switching from immediate release to extended-release product, use same total daily dose. The immediate and extended release products may not give same clinical response on mg:mg basis; monitor response and side effects when interchanging between metoprolol products. |
| | Concomitant amiodarone, digoxin, disopyramide, or non-dihydropyridine calcium channel blockers may increase the risk of bradycardia. |
| | Monitor closely for HF exacerbation and hypotension when titrating dose. |
| Counseling | Do not abruptly discontinue without physician's advice. |
| | Take with food or directly after eating. |
| | Extended-release tablets may be broken in half, but do not chew or crush. |

^{*}Refer to prescribing information for more complete information.

Sources:

- 1. American College of Cardiology (ACC), American Heart Association (AHA), and the European Society of Cardiology (ESC). ACC/AHA/ESC 2006 Guidelines for the Management of Patients With Atrial Fibrillation. Washington, DC: American College of Cardiology.
- 2. Heart Rhythm Society. *AF360 Pocket Guide: Practical Rate and Rhythm Management of Atrial Fibrillation*. 2010, Washington, DC: Heart Rhythm Society.
- 3. Tarascon Pocket Pharmacopoeia®2012.

[†]Dosages given in the table may differ from those recommended by the manufacturers.