

**Initial REMS Approval: 09/2009**  
**Most Recent Modification: 05/2015**

**NDA 21-454 TESTIM<sup>®</sup> (testosterone gel) CIII**  
**and Authorized Generic (testosterone gel) CIII**

**Class of Drug: Androgen**

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## **RISK EVALUATION AND MITIGATION STRATEGY (REMS)**

### **I. GOAL:**

The goal of this REMS is to inform patients about the serious risks associated with the use of testosterone gel.

### **II. REMS ELEMENTS**

#### **A. Medication Guide**

A Medication Guide will be dispensed with each testosterone gel prescription in accordance with 21 CFR 208.24.

The Medication Guides are part of the REMS.

#### **B. Timetable for Submission of Assessments**

Auxilium Pharmaceuticals will submit REMS Assessment to FDA for testosterone gel by 18 months, 3 years, and 7 years from the date of initial approval (09/18/2009) of the TESTIM<sup>®</sup> (testosterone gel) REMS.

To facilitate inclusion of as much information as possible while allowing reasonable time to prepare the submission, the reporting interval covered by each assessment should conclude no earlier than 60 days before the submission date for that assessment. Auxilium Pharmaceuticals will submit each assessment so that it will be received by the FDA on or before the due date.