

## RxAdmix - Key Features that meet the California Board of Pharmacy Regulations

- 1) Provides a Master Formula Record that includes the following elements:
  - a. Active ingredients used
  - b. Equipment to be used
  - c. Expiration dating requirements
  - d. Inactive ingredients used
  - e. Process and/or procedure used to prepare the drug
  - f. Quality reviews required at each step in the preparation of the drug
  - g. Post-compounding process or procedures if required
- 2) Provides an online fill-able Pharmacy Self-Assessment Form
- 3) Provides for the record keeping of the following items:
  - a. The master formula record
  - b. The date the drug product was compounded
  - c. The identity of the person who compounded the drug
  - d. The identity of the pharmacist reviewing the final drug product
  - e. The quantity of each component principal active ingredient(s) and diluent(s)
  - f. The expiration date of the final compounded drug product
  - g. Unique "Pharmacy Reference Number" called the Compound's Lot Number for each unique compound
  - h. The manufacturer, lot number and expiration date of each principal active ingredient(s) and diluent(s)
  - i. The quantity or amount of drug product compounded
- 4) Identifies storage requirements for each unique compound
- 5) Identifies the equipment used to prepare each unique compound.
- 6) Provides label printing for the compounds that meet the pharmacy board requirements and ISMP recommendations. The labels contain:
  - a. The generic names of each principal active ingredient(s) and diluent(s)
  - b. "Pharmacy Reference Number" called the Compound's Lot Number
  - c. Concentration of Strength
  - d. Volume or weight
  - e. Expiration date
  - f. Including warning labels and icons such as: "Chemotherapy-Dispose of properly" for all cytotoxic agents
  - g. Telephone number of the pharmacy
  - h. Instructions for storage and handling