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