

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

- 1) Provides for a Master Formula Record. It includes the following elements:
  - a. Active ingredients used
  - b. Inactive ingredients used
  - c. Process and/or procedure used to prepare the drug.
  - d. Quality reviews required at each step in the preparation of the drug.
  - e. Post-compounding process or procedures if required.
  - f. Expiration dating requirements
  - g. Storage requirements
- 2) Provides an online fill-able Pharmacy Self-Assessment Form.
- 3) Provides for the record keeping of the:
  - a. Identity of the person who compounded the drug.
  - b. Identity of the pharmacist reviewing the final drug product.
  - c. NDC # of each principal active ingredient(s) and diluent(s).
  - d. The lot number and expiration date of each principal active ingredient(s) and diluent(s).
  - e. The quantity of each component principal active ingredient(s) and diluent(s).
  - f. Unique "Pharmacy Reference Number" called the Compound's Lot Number for each unique compound.
  - g. The date the drug product was compounded
  - h. The expiration date of the final compounded drug product.
  - i. The quantity or amount of drug product compounded.
- 4) Identifies storage requirements for each unique compound.
- 5) Identifies the expiration date for each unique compound.
- 6) Identifies the equipment to be used to prepare each unique compound.
- 7) Provides for the printing of labels for the compounds meeting pharmacy board requirements and ISMP recommendation. The labels contain at least:
  - 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;
    - i. "Chemotherapy-Dispose of Properly" for all cytotoxic agents.