RxScan
Preventing Medication Errors

Perfection Is Your Goal, We Help You Get There™

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.

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