DRUG UTILIZATION REVIEW (DUR)



DUR

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DUR: Legal Basis

- Federal law requires each state Medicaid program to develop a a DUR program, including establishing a DUR Board. DUR activities must include:
 - ✓ Retrospective DUR
 - √ Prospective DUR
 - ✓ Educational activities and interventions.
- Omnibus Budget Reconciliation Act of 1990 (OBRA 90), 42 CFR Part 456 Utilization Control, subpart K— Drug Use Program and Electronic Claims Management System for Outpatient Drug Claims.
- Program must be effective by January 1993

DUR Board Responsibilities

- Review and make recommendations about predetermined standards for retrospective and prospective DUR criteria.
- Evaluate the use of predetermined standards concerning modification or elimination of existing standards or the addition of new ones.

DUR BOARD MEMBERSHIP

- At lease one-third, but not more than 51 percent must be physicians.
- At least one-third must be pharmacists.
- The physicians and pharmacists must be actively practicing and licensed.

What is DUR?

DUR is Drug Utilization Review.
 There are three required activities for DUR: Prospective and Retrospective as well as Educational Programs and Interventions.

DUR: Purpose

"To improve the quality of pharmaceutical care by insuring prescriptions are appropriate, medically necessary, and that they are not likely to result in adverse medical results."

Prospective DUR

- Review of drug claims that have been verified as payable through the point of sale system.
- Since this review occurs prior to the dispensing of the drug, the primary audience for follow up is pharmacists.
- Some predetermined standards include:
 - ✓ Early refill.
 - ✓ Drug-Drug Interaction.
 - ✓ Overutilization.

Prospective DUR

- ✓ Improve the quality and cost effectiveness of drug use.
- ✓ Verify prescriptions dispensed are appropriate, medically necessary and not likely to result in adverse drug events.
- √ Verification occurs before the drug is dispensed.
- ✓ Allows pharmacist an opportunity to perform patient counseling.

- PROSPECTIVE DUR: Specific projects
 - ✓ Review of pharmacy use of alerts with targeted letter interventions to outliers.*
 - ✓ Preparation of drug list for point of sale (POS) monitoring to avoid adverse drug reactions (ADE) – Drug/Drug; Drug/Diagnosis; etc.*
 - *Educational programs and interventions developed as a result of prospective DUR reviews.

Retrospective DUR

- Monthly review of drug claims data for potential drug use problems.
- Since this review occurs after the drugs have been dispensed, the primary audience for follow up is the prescriber.
- Some predetermined standards include:
 - ✓ Overutilization
 - √ Therapeutic duplication
 - ✓ Additive Toxicity

Retrospective DUR

- Primary audience is the prescriber
- Also provides for additional monitoring of appropriate prescribing
- Detect fraud, abuse, overuse or medically unnecessary use of medication.
- Recipient and prescriber targets are identified for educational intervention(s).

- Retrospective DUR projects:
 - Analysis of anti-emetic drug use
 - Off-label use of epileptic drugs
 - Asthma interventions
 - Post-MI intervention

These were educational programs and interventions developed as a result of retrospective DUR reviews.

DUR Board Responsibilities

- Educational programs including interventions
 - ✓ Identify and develop educational topics to improve prescribing and dispensing practices.
 - ✓ Recommend appropriate interventions based on in-depth review of claims review of claims data
 - ✓ Periodically re-evaluate and modify interventions, if necessary

Educational Programs and Interventions

 Based on findings from reviews of Prospective and Retrospective DUR, the Board identifies and develops educational topics to improve prescribing and dispensing practices.

- Recipient Lock-in
 - Coordinate the provision of health care services for recipients who abuse or misuse Medicaid benefits
 - Improve the quality of care for the recipient and reduce unnecessary physician and pharmacy utilization
 - Allow recipient reasonable access to necessary Medicaid services

Recipient Lock-in: how it works

- Candidates for Lock-In come from referrals from retroDUR, physicians, pharmacists, and other health care providers
- Decision Support Tool is an automated process for identifying recipients for potential lock-in.
- 6 months of pharmacy claims and diagnosis data reviewed by pharmacist
- APS provides recommendations including:
 - Alert letter to physicians
 - Warning letter to recipient
 - Lock-In

Recipient Lock-in: how it works (con't)

- If the recipient is recommended for lock-in and Division of Health Care Financing (DHCF) agrees, then the recipient:
 - Receives letter of intent to lock-in
 - Letter explains restrictions to be applied
 - How to designate a physician and pharmacy
 - How to request a hearing within 15 days
 - If recipient fails to designate providers the RLP may assign providers based on claims history

Recipient Lock-in: how it works (con't)

- APS oversees coordination of care for 24 months of lock-in enrollment
 - Monitor claims payment
 - Track physicians, pharmacies and referrals
 - Respond to provider and recipient inquiries
 - One month before recipient's scheduled release paid and denied pharmacy claims are reviewed for compliance with guidelines

DRUG UTILIZATION REVIEW Annual Objectives

• SFY 2006

- On the basis of paid claims analysis choose and complete two interventions to improve utilization of pharmacy benefit
- Review current alerts that are monitored on a quarterly basis and recommend changes to improve monitoring results
- Choose two newsletter topics based on subjects chosen by participating pharmacists

DRUG UTILIZATION REVIEW Staff

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