Revision: HCFA-AT-80-38 (BPP)

May 22, 1980

State/Territory: <u>NEVADA</u>

Citation

4.25 <u>Program for Licensing Administrators of Nursing Homes</u>

42 CFR 431.702 AT-78-90

The State has a program that, except with respect to Christian Science sanatoria, meets the requirements of 42 CFR Part 431, Subpart N, for the licensing of nursing home administrators.

OMB No.: 0938-0193

TN No. <u>74-3</u> Approval Date: <u>5/15/74</u> Effective Date: <u>12/1/73</u>

Supersedes TN No. \_\_\_\_

Revision: HCFA-PM-93-3

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Citation

1927(g)

4.2 Drug Utilization Review Program

(MB)

42 CFR 456.700

- A.1. The Medicaid agency meets the requirements of Section 1927(g) of the Act for a drug use review (DUR) program for outpatient drug claims.
  - 2. The DUR program assures that the prescriptions for outpatient drugs are:
    - Appropriate
    - Medically necessary
    - Are not likely to result in adverse medical results
- B. The DUR program is designed to educate physicians and pharmacists to identify and reduce the frequency of patterns of fraud, abuse, gross overuse, or inappropriate or medically unnecessary care among physicians, pharmacists, and patients or associated with specific drugs as well as:
  - Potential and actual adverse drug reactions
  - Therapeutic appropriateness
  - Overutilization and under utilization
  - Appropriate use of generic products
  - Therapeutic duplication
  - Drug disease contraindications
  - Drug-drug interactions
  - Incorrect drug dosage or duration of drug treatment
  - Drug-allergy interactions
  - Clinical abuse/misuse

TN No. 93-10 Approval Date: May 7, 1993 Effective Date: January 1, 1993

74a

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1927(g)(1)(B)42 CFR 456.703 (d) and (f)

- C. The DUR program shall assess data use against predetermined standards whose source materials for their development are consistent with peer-reviewed medical literature which has been critically reviewed by unbiased independent experts and the following compendia:
  - American Hospital Formulary Service Drug Information
  - United States Pharmacopia-Drug Information
  - American Medical Association Drug Evaluations

1927(g)(1)(D)42 CFR 456.703(b)

DUR is not required for drugs dispensed to residents of nursing D. facilities that are in compliance with drug regiment review procedures set forth in 42 CFR 483.60. The State has never-theless chosen to include nursing home drugs in:

> Prospective DUR X Retrospective DUR

1927(g)(2)(A)42 CFR 456.705(b) E.1. The DUR program includes prospective review of drug therapy at the point of sale or point of distribution before each prescription is filled or delivered to the Medicaid recipient.

1927(g)(2)(A)(i)42 CFR 456.705(b), (1)-(7))

- 2. Prospective DUR includes screening each prescription filled or delivered to an individual receiving benefits for potential drug therapy problems due to:
  - Therapeutic duplication
  - Drug-disease contraindications
  - Drug-drug interactions
  - Drug-interactions with non-prescription or over-the-counter

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- Incorrect drug dosage or duration of drug treatment
- Drug allergy interactions
- Clinical abuse/misuse

Approval Date: May 7, 1993

Revision: HCFA-PM-93-3 (MB) March 1993 State/Territory: **NEVADA** 3. 1927(g)(2)(A)(ii)Prospective DUR includes counseling for Medicaid 42 CFR 456.705(c) recipients based on standards established by state law and maintenance of patient profiles. and (d) F.1 The DUR program includes retrospective DUR through its 1927(g)(2)(B)42 CFR 456.709(a) mechanized drug claims processing and information retrieval system or otherwise which undertakes ongoing periodic examination of claims data and other records to identify: Patterns of fraud and abuse Gross overuse Inappropriate or medically unnecessary care among physicians, pharmacists, Medicaid recipients, or associated with specific drugs or groups of drugs. F.2 1927(g)(2)(C)The DUR program assesses data on drug use against 42 CFR 456.709(b) explicit predetermined standards including but not limited to monitoring for: Therapeutic appropriateness Overutilization and underutilization Appropriate use of generic products Therapeutic duplication Drug-disease contraindications Drug-drug interactions Incorrect drug dosage/duration of drug treatment Clinical abuse/misuse 1927(g)(2)(D)3. The DUR program through its State DUR Board, using data provided by the Board, provides for active and ongoing 42 CFR 456.711 educational outreach programs to educate practitioners on common drug therapy problems to improve prescribing and dispensing practices. G.1 The DUR program has established a State DUR Board 1927(g)(3)(A)42 CFR 456.716(a) either: Directly, or X Under contract with a private organization TN No. 93-10 Approval Date: May 7, 1993 Effective Date: January 1, 1993 Supersedes

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**NEVADA** 

(MB)

1927(g)(3)(B)42 CFR 456.716 (A) and (B)

- 2. The DUR Board membership includes health professionals (one-third licensed actively practicing pharmacists and onethird but no more than 51 percent licensed and actively practicing physicians) with knowledge and experience in one or more of the following:
  - Clinically appropriate prescribing of covered outpatient
  - Clinically appropriate dispensing and monitoring of covered outpatient drugs
  - Drug use review, evaluation and intervention
  - Medical quality assurance

1927(g)(3)(C)42 CFR 456.716(d)

- 3. The activities of the DUR Board include:
  - Retrospective DUR;
  - Application of Standards as defined in section 1927(g)(2)(C); and
  - Ongoing interventions for physicians and pharmacists targeted toward therapy problems or individuals identified in the source of retrospective DUR.

1927(g)(3)(C)42 CFR 456.711 (a)-(d)

- G.4 The interventions include in appropriate instances:
  - Information dissemination
  - Written, oral, and electronic reminders
  - Face-to-face discussions
  - Intensified monitoring/review of prescribers/dispensers

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H.

I.1

J.

1927(g)(D) 42 CFR 456.712 (A) and (B) The State assures that it will prepare and submit an annual report to the secretary, which incorporates a report from the State DUR Board and that the State will adhere to the plans, steps, procedures as described in the report.

1927(h)(1) 42 CFR 456.722 The State establishes, as its principal means of processing claims for covered outpatient drugs under this title, a pointof-sale electronic claims management system to perform on-line:

- Real time eligibility verification
- Claims data capture
- Adjudication of claims
- Assistance to pharmacists, etc., applying for and receiving payment

1927(g)(2)(A)(i) 42 CFR 456.705(b) 2. Prospective DUR is performed using an electronic point of sale drug claims processing system.

1927(j)(2) 42 CFR 456.703(c) Hospitals which dispense covered outpatient drugs are exempted from the drug utilization review requirements of this section when facilities use drug formulary systems and bill the Medicaid program no more than the hospital's purchasing cost for such covered outpatient drugs. The hospitals will provide documentation to the State to allow the State to make such exemptions.