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State/Territory Name: Oklahoma

State Plan Amendment (SPA)#: 20-0038

This file contains the following documents in the order listed

- 1) Approval Letter
- 2) CMS 179 Form
- 3) Approved SPA Pages

DEPARTMENT OF HEALTH & HUMAN SERVICES Centers for Medicare & Medicaid Services 7500 Security Boulevard, Mail Stop S2-14-26 Baltimore, Maryland 21244-1850



<u>Center for Medicaid and CHIP Services</u> Disabled and Elderly Health Programs Group

November 23, 2020

Ms. Melody Anthony State Medicaid Director Oklahoma Health Care Authority 4345 N. Lincoln Blvd. Oklahoma City, OK 73105

Dear Ms. Anthony,

The CMS Division of Pharmacy team has reviewed Oklahoma's State Plan Amendment (SPA) 20-0038 received in the CMS Medicaid & CHIP Operations Group on November 4, 2020. This SPA proposes to update the standard for retrospective drug utilization reviews (DUR) in accordance with section 1004 of the Substance Use-Disorder Prevention that promotes Opioid Recovery and Treatment (SUPPORT) for Patients and Communities Act (P.L. 115-271).

Based on the information provided and consistent with the regulations at 42 CFR 430.20, we are pleased to inform you that OK-20-0038 is approved with an effective date of October 1, 2020. We are attaching a copy of the signed CMS-179 form, as well as the pages approved for incorporation into Oklahoma's state plan.

If you have any questions regarding this request, please contact Michael Forman at 410-786-2666 or michael.forman@cms.hhs.gov.

Sincerely,

John M. Coster, Ph.D., R.Ph. Director Division of Pharmacy

cc: Amanda Salisbury, Oklahoma Health Care Authority
Traylor Rains, Oklahoma Health Care Authority
Sandra Puebla, Oklahoma Health Care Authority
Deborah Read, CMS, Medicaid & CHIP Operations Group

CENTERS FOR MEDICARE & MEDICAID SERVICES		OIVID INU. 0930-0193
TRANSMITTAL AND NOTICE OF APPROVAL OF STATE PLAN MATERIAL	1. TRANSMITTAL NUMBER	2. STATE
FOR: CENTERS FOR MEDICARE & MEDICAID SERVICES	3. PROGRAM IDENTIFICATION: TITLE > SECURITY ACT (MEDICAID)	(IX OF THE SOCIAL
TO: REGIONAL ADMINISTRATOR CENTERS FOR MEDICARE & MEDICAID SERVICES DEPARTMENT OF HEALTH AND HUMAN SERVICES	4. PROPOSED EFFECTIVE DATE	
5. TYPE OF PLAN MATERIAL (Check One)	•	
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COMPLETE BLOCKS 6 THRU 10 IF THIS IS AN AMENI		mendment)
6. FEDERAL STATUTE/REGULATION CITATION	7. FEDERAL BUDGET IMPACT a. FFY\$ b. FFY\$	
8. PAGE NUMBER OF THE PLAN SECTION OR ATTACHMENT	9. PAGE NUMBER OF THE SUPERSI OR ATTACHMENT (If Applicable)	EDED PLAN SECTION
10. SUBJECT OF AMENDMENT		
11. GOVERNOR'S REVIEW (Check One)		
☐ GOVERNOR'S OFFICE REPORTED NO COMMENT☐ COMMENTS OF GOVERNOR'S OFFICE ENCLOSED☐ NO REPLY RECEIVED WITHIN 45 DAYS OF SUBMITTAL	OTHER, AS SPECIFIED	
12. SIGNATURE OF STATE AGENCY OFFICIAL	6. RETURN TO	
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19. EFFECTIVE DATE OF APPROVED MATERIAL October 1, 2020	D. SIGNATURE OF REGIONAL OFFICIA	AL .
21. TYPED NAME John M. Coster, Ph.D., R.Ph.	2. TITLE Director, Division of Pha	rmacy
23. REMARKS		

Revision:	HCFA-PM-
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(MB)

State/Territory: OKLAHOMA

Citation

1927(g)(1)(D) 42 CFR 456.703(b) DUR is not required for drugs dispensed to residents of nursing facilities that are in compliance with drug regimen review procedures set forth in 42 CFR 483.60. The State has never-the-less chose to include nursing home drugs in:

X 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 interaction
 - Drug interactions with no-prescription or over-the-counter drugs
 - Incorrect drug dosage or duration of drug treatment
 - Drug-allergy interactions
 - Clinical abuse/misuse

At the option of the State, the screenings also include review for:

- High drug dosages
- Drug age precaution
- Drug-pregnancy
- Ingredient duplication

1927(g)(2)(A)(ii) 42 CFR 456.705(c) and d

2. Prospective DUR includes counseling for Medicaid recipients based on standards established by State law and maintenance of patient profiles.

1927(g)(2)(B) 42 CFR 456.709(a)

- F.1. The DUR program includes retrospective DUR through its 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
 - Excessive utilization
 - Inappropriate or medically unnecessary care or prescribing or billing practices that indicate abuse or excessive utilization among physicians, pharmacists, Medicaid members, or associated with specific drugs or groups of drugs.

TN #: 20-0038 Effective Date: 10/01/2020

Supersedes TN #: 19-0040 Approval Date: November 23, 2020