

Table of Contents

State/Territory Name: Tennessee

State Plan Amendment (SPA) #: 19-0005

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



Center for Medicaid and CHIP Services

Disabled and Elderly Health Programs Group

March 13, 2020

Mr. Stephen M. Smith
Director
Division of TennCare
310 Great Circle Road
Nashville, Tennessee 37243

Dear Mr. Smith:

The CMS Division of Pharmacy team has reviewed Tennessee's State Plan Amendment (SPA) 19-0005 received in the CMS Division of Program Operations on December 23, 2019. This SPA proposes to allow the state to comply with the Medicaid Drug Utilization Review (DUR) provisions included in 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 SPA 19-0005 is approved with an effective date of October 1, 2019. A copy of the signed CMS-179 form, as well as the pages approved for incorporation into Tennessee's state plan, will be forwarded by the CMS Division of Program Operations.

If you have any questions regarding this request, please contact Lisa Shochet at (410) 786-5445 or Lisa.Shochet@cms.hhs.gov.

Sincerely,

/s/

Cynthia R. Denemark, R.Ph.
Deputy Director
Division of Pharmacy
DEHPG/CMCS/CMS

cc: George Woods, Division of TennCare
James G. Scott, Division Director, CMS Division of Program Operations
Tandra Hodges, CMS Division of Program Operations

Revision: HCFA-PM- (MB)

OMB No.

State/Territory: TennesseeCitation

1902(a)(85) and
Section 1004 of the
Substance Use-
Disorder Prevention that
Promotes Opioid
Recovery and
Treatment for Patients
and Communities
Act (SUPPORT Act)

K. 1. Claims Review Limitations:

Prospective safety edits on opioid prescriptions to address days' supply, early refills, duplicate fills, and quantity limitations for clinical appropriateness.

Prospective safety edits on maximum daily morphine milligram equivalents (MME) on opioids prescriptions to limit the daily morphine milligram equivalent (as recommended by clinical guidelines).

Retrospective reviews on opioid prescriptions exceeding these above limitations on an ongoing basis.

Retrospective reviews on concurrent utilization of opioids and benzodiazepines as well as opioids and antipsychotics on an ongoing periodic basis.

2. Programs to Monitor Antipsychotic Medications to children:

Antipsychotic agents are reviewed for appropriateness for all children, including foster children, based on approved indications and clinical guidelines.

3. Fraud and Abuse Identification:

The DUR board participates in established initiatives (including the State's pharmacy lock-in program) that identify potential fraud or abuse of controlled substances by enrolled individuals, health care providers, and pharmacies.

TN No.: 19-0005
Supersedes
TN No.: New

Approval Date: 03/13/2020Effective Date: 10/1/19