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**State/Territory Name: Rhode Island** 

State Plan Amendment (SPA) #:19-015

This file contains the following documents in the order listed:

- 1) Approval Letter
- 2) 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

February 6, 2020

Womazetta Jones, Secretary Executive Office of Health and Human Services State of Rhode Island 3 West Road Cranston, Rhode Island 02920

Dear Ms. Jones:

The CMS Division of Pharmacy team has reviewed Rhode Island State Plan Amendment (SPA) 19-0015 received in the Boston Regional Operations Group on December 19, 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 19-0015 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 Rhode Island's state plan will be forwarded by the Boston Regional Operations Group.

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: James G. Scott, Division Director, CMS Division of Program Operations Lynn DelVecchio, Boston Regional Operations Group Joyce Butterworth, Boston Regional Operations Group Revision: HCFA-PM- (MB)

State/Territory: Rhode Island

#### <u>Citation</u>

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)

#### **Claim 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

## 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

**Fraud and abuse identification:** The Surveillance Utilization Review (SUR) team has established a process that identifies potential fraud or abuse of controlled substances by enrolled individuals, health care providers and pharmacies.

TN:<u>19-0015</u>
Supersedes Approved: <u>02/06/2020</u> Effective: <u>October 1, 2019</u>

TN: NEW