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State/Territory Name: Arkansas

State Plan Amendment (SPA) #: 23-0009

This file contains the following documents in the order listed:

Approval Letter
CMS Form 179
Approved SPA Pages

DEPARTMENT OF HEALTH & HUMAN SERVICES Centers for Medicare & Medicaid Services 7500 Security Boulevard, Mail Stop S2-26-12 Baltimore, Maryland 21244-1850



May 10, 2023

Janet Mann Director of Health and Medicaid Director Arkansas Department of Human Services 112 West 8th Street, Slot S401 Little Rock, AR 72201-4608

Re: Arkansas State Plan Amendment (SPA) AR-23-0009

Dear Director Mann:

The Centers for Medicare & Medicaid Services (CMS) reviewed the proposed Medicaid State Plan Amendment (SPA) and accompanying section 1135 waivers submitted on March 31, 2023 under transmittal number (TN) AR-23-0009. This amendment proposes to add coverage of routine patient costs associated with participation in qualifying clinical trials.

Pursuant to section 1135(b)(5) of the Social Security Act (Act), for the period of the public health emergency, CMS is modifying the requirement at 42 C.F.R. §430.20 that the state submit SPAs related to the COVID-19 public health emergency by the final day of the quarter, to obtain a SPA effective date during the quarter, enabling SPAs submitted after the last day of the quarter to have an effective date in a previous quarter, but no earlier than the effective date of the public health emergency.

The State of Arkansas also requested a waiver to modify the public notice requirements applicable to the SPA submission process. Pursuant to section 1135(b)(5) of the Act, CMS is modifying public notice requirements applicable to the SPA submission process. Public notice for SPAs is required under 42 C.F.R. §447.205 for changes in statewide methods and standards for setting Medicaid payment rates, 42 C.F.R. §447.57 for changes to premiums and cost sharing, and 42 C.F.R. §440.386 for changes to Alternative Benefit Plans (ABPs). Pursuant to section 1135(b)(5) of the Act, CMS is approving the state's request to modify these notice requirements otherwise applicable to SPA submissions.

CMS conducted our review of your submittal according to statutory requirements in Title XIX of the Act and implementing regulations. This letter is to inform you that Arkansas' Medicaid SPA Transmittal Number AR-23-0009 is approved effective January 1, 2022.

Please contact Lee Herko at 570-230-4048 or by email at <u>Lee.Herko@cms.hhs.gov</u> if you have any questions about this approval.

Sincerely,

Alissa M. Deboy -S

Digitally signed by Alissa M. Deboy -S Date: 2023.05.10 08 29:33 -04'00'

Alissa Mooney DeBoy On Behalf of Anne Marie Costello, Deputy Director Center for Medicaid and CHIP Services

Enclosures

DEPARTMENT OF HEALTH AND HUMAN SERVICES	FORM APPROVED OMB No 0938-0193
CENTERS FOR MEDICARE & MEDICAID SERVICES	1. TRANSMITTAL NUMBER 2. STATE
TRANSMITTAL AND NOTICE OF APPROVAL OF	2 3 - 0 0 0 9 A R
STATE PLAN MATERIAL	3. PROGRAM IDENTIFICATION: TITLE OF THE SOCIAL
FOR: CENTERS FOR MEDICARE & MEDICAID SERVICES	SECURITY ACT XIX XXI
TO: CENTER DIRECTOR	4. PROPOSED EFFECTIVE DATE
CENTERS FOR MEDICAID & CHIP SERVICES DEPARTMENT OF HEALTH AND HUMAN SERVICES	1/1/2022
5. FEDERAL STATUTE/REGULATION CITATION	6. FEDERAL BUDGET IMPACT (Amounts in WHOLE dollars)
Section 1905(gg)(1) – 1905(gg)(3) of the Social Security Act	a FFY\$_0
SMD # 21-005	b. FFY2024\$_0
7. PAGE NUMBER OF THE PLAN SECTION OR ATTACHMENT	8. PAGE NUMBER OF THE SUPERSEDED PLAN SECTION OR ATTACHMENT (If Applicable)
Attachment 3.1-A Page 13	
Attachment 3.1-B Page 12	Attachment 3.1-A Page 13
	Attachment 3.1-B Page 12
9. SUBJECT OF AMENDMENT	
Clinical Trials Attestation	
10. GOVERNOR'S REVIEW (Check One)	
GOVERNOR'S OFFICE REPORTED NO COMMENT OTHER, AS SPECIFIED:	
NO REPLY RECEIVED WITHIN 45 DAYS OF SUBMITTAL	
	15. RETURN TO
CIAL	15. RETORN TO
	Office of Rules Promulgation
	PO Box 1437, Slot S295
13. TITLE	Little Rock, AR 72203-1437
Director, Division of Medical Services	Attn: Mac Golden
14. DATE SUBMITTED	
3-31-2023	ISE ONLY
	17. DATE APPROVED
March 31, 2023	May 10, 2023
PLAN APPROVED - OI	
18. EFFECTIVE DATE OF APPROVED MATERIAL	19. SIGNATURE OF APPROVING STEAL M. Deboy -S
January 1, 2022	Deboy -S Date: 2023.05.10 08 30:19 -04'00'
20. TYPED NAME OF APPROVING OFFICIAL	21. TITLE OF APPROVING OFFICIAL
Alissa Mooney DeBoy	On Behalf of Anne Marie Costello, Deputy Director, CMCS
22. REMARKS	
Pen and Ink Request Approved Boxes 5 and 8: Pen and ink requests by State on 5/1/2023 to correct Box approved by State on 5/08/2023.	
by State on 5/1/2023 to correct Box approved by State on 5/08/2023. 14: Date Submitted 03/31/2023.	

ATTACHMENT 3.1-A

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State/Territory: Arkansas

AMOUNT, DURATION AND SCOPE OF SERVICES PROVIDED

CATEGORICALLY NEEDY GROUP(S)

30. Coverage of Routine Patient Cost in Qualifying Clinical Trials

*The state needs to check each assurance below.

Provided: _01/01/2022

I. General Assurances:

Routine Patient Cost – Section 1905(gg)(1)

X ___ Coverage of routine patient cost for items and services as defined in section 1905(gg)(1) that are furnished in connection with participation in a qualified clinical trial.

Qualifying Clinical Trial – Section 1905(gg)(2)

_X_A qualified clinical trial is a clinical trial that meets the definition at section 1905(gg)(2).

Coverage Determination – Section 1905(gg)(3)

X A determination with respect to coverage for an individual participating in a qualified clinical trial will be made in accordance with section 1905(gg)(3).

PRA Disclosure Statement - This information is being collected to assist the Centers for Medicare & Medicaid Services in implementing Section 210 of the Consolidated Appropriations Act of 2021 amending section 1905(a) of the Social Security Act (the Act), by adding a new mandatory benefit at section 1905(a)(30). Section 210 mandates coverage of routine patient services and costs furnished in connection with participation by Medicaid beneficiaries in qualifying clinical trials effective January 1, 2022. Section 210 also amended sections 1902(a)(10)(A) and 1937(b)(5) of the Act to make coverage of this new benefit mandatory under the state plan and any benchmark or benchmark equivalent coverage (also referred to as alternative benefit plans, or ABPs). Under the Privacy Act of 1974 any personally identifying information obtained will be kept private to the extent of the law. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unlessit displays a currently valid Office of Management and Budget (OMB) control number. The OMB control number for this project is 0938-1148 (CMS-10398 #74). Public burden for all of the collection of information requirements under this control number is estimated to take about 56 hours per response. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to CMS, 7500 SecurityBoulevard, Attn: Paperwork Reduction Act Reports Clearance Officer, Mail Stop C4-26-05, Baltimore, Maryland 21244-1850.

ATTACHMENT 3.1-B

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State/Territory: Arkansas

AMOUNT, DURATION AND SCOPE OF SERVICES PROVIDED

MEDICALLY NEEDY GROUP(S)

30. Coverage of Routine Patient Cost in Qualifying Clinical Trials

*The state needs to check each assurance below.

Provided: X

I. General Assurances:

Routine Patient Cost – Section 1905(gg)(1)

 $X_Coverage of routine patient cost for items and services as defined in section 1905(gg)(1) that are furnished in connection with participation in a qualified clinical trial.$

Qualifying Clinical Trial – Section 1905(gg)(2)

_X__A qualified clinical trial is a clinical trial that meets the definition at section 1905(gg)(2).

Coverage Determination – Section 1905(gg)(3)

 X_A determination with respect to coverage for an individual participating in a qualified clinical trial will be made in accordance with section 1905(gg)(3).

PRA Disclosure Statement - This information is being collected to assist the Centers for Medicare & Medicaid Services in implementing Section 210 of the Consolidated Appropriations Act of 2021 amending section 1905(a) of the Social Security Act (the Act), by adding a new mandatory benefit at section 1905(a)(30). Section 210 mandates coverage of routine patient services and costs furnished in connection with participation by Medicaid beneficiaries in qualifying clinical trials effective January 1, 2022. Section 210 also amended sections 1902(a)(10)(A) and 1937(b)(5) of the Act to make coverage of this new benefit mandatory under the state plan and any benchmark or benchmark equivalent coverage (also referred to as alternative benefit plans, or ABPs). Under the Privacy Act of 1974 any personally identifying information obtained will be kept private to the extent of the law. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unlessit displays a currently valid Office of Management and Budget (OMB) control number. The OMB control number for this project is 0938-1148 (CMS-10398 #74). Public burden for all of the collection of information requirements under this control number is estimated to take about 56 hours per response. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to CMS, 7500 SecurityBoulevard, Attn: Paperwork Reduction Act Reports Clearance Officer, Mail Stop C4-26-05, Baltimore, Maryland 21244-1850.

TN: <u>23-0009</u> Supersedes TN: New Page Approval Date: <u>05/10/2023</u> Effective Date: <u>01/01/2022</u>