### **Attachment 3.1A: Freestanding Birth Center Services**

TN <u>15-0038</u>

Provided:	☐ No limitations	With limitations	☐None licensed or approved		
Stays and the service practific immediate.  The From with we that all the service practifications are serviced as the service practification of the service	ne services rendered for lates rendered during input tioner from the time of the diate postpartum period.  SBC shall be located with which the FSBC shall materials.	abor and delivery are very ient hospital stays. Service pregnant woman's admethin a ground travel time of the contractual relation caesarian delivery to begin	FSBC) are typically less than 24 hours y limited in comparison to delivery es shall be provided by the attending hission through the birth and the distance from a general acute care hospital onship, including a transfer agreement, in within 30 minutes of the decision a		
	nsed or Otherwise Sta	te-Recognized covered	professionals providing services in		
Provided:	☐ No limitations		lease describe below)		
☐ Not A	Not Applicable (there are no licensed or State approved Freestanding Birth Centers)				
	ibe any limitations: standing birthing center s	staff shall not administer §	general or epidural anesthesia services.		
Please check	all that apply:				
	<del>-</del>	<u>=</u>	scribed in another benefit category ns and certified nurse midwives).		
care in a free are otherwis	estanding birth center	within the scope of pra R 440.60 (e.g., lay mid	I, labor and delivery, or postpartum ctice under State law whose services wives, certified professional midwives		
— · ·	•	ssionals licensed or otherces (e.g., doulas, lactat	erwise recognized by the State to ion consultant, etc.).*		
providing bir	(c) above, please list a th center services: sed midwives	nd identify below each	type of professional who will be		

State/Territory: <u>Louisiana</u>

# 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: With limitations

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).

#### **Limitations:**

Routine patient costs in qualified clinical trials do not include the following:

- 1. Any investigational item or service that is the subject of the qualifying clinical trial and is not otherwise covered outside of the clinical trial under the State Plan; and
- 2. Any item or service that is provided to the beneficiary solely to satisfy data collection and analysis needs for the qualifying clinical trial and not used in the direct clinical management of the beneficiary and that is not otherwise covered under the State Plan.

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 notconduct or sponsor, and a person is not required to respond to, a collection of information unless it 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: 22-0012 Approval Date: Effective Date: January 1, 20	2022
--	------

Supersedes TN: New Page

### **Attachment 3.1B: Freestanding Birth Center Services**

TN New Page

29. (i) Licensed or Otherwise State-Approved Freestanding Birth Centers
Provided: ☐ No limitations ☐ With limitations ☐ None licensed or approved
Stays for delivery at the free-standing birthing centers (FSBC) are typically less than 24 hours and the services rendered for labor and delivery are very limited in comparison to delivery services rendered during inpatient hospital stays. Services shall be provided by the attending practitioner from the time of the pregnant woman's admission through the birth and the immediate postpartum period.
The FSBC shall be located within a ground travel time distance from a general acute care hospital with which the FSBC shall maintain a contractual relationship, including a transfer agreement, that allows for an emergency caesarian delivery to begin within 30 minutes of the decision a caesarian delivery is necessary.
29. (ii) Licensed or Otherwise State-Recognized covered professionals providing services in the Freestanding Birth Center
Provided: ☐ No limitations ☐ with limitations (please describe below)
Not Applicable (there are no licensed or State approved Freestanding Birth Centers)
Please describe any limitations:  Free-standing birthing center staff shall not administer general or epidural anesthesia services.
Please check all that apply:
(a) Practitioners furnishing mandatory services described in another benefit category and otherwise covered under the State plan (i.e., physicians and certified nurse midwives).
(b) Other licensed practitioners furnishing prenatal, labor and delivery, or postpartum care in a freestanding birth center within the scope of practice under State law whose services are otherwise covered under 42 CFR 440.60 (e.g., lay midwives, certified professional midwives (CPMs), and any other type of licensed midwife). *
(c) Other health care professionals licensed or otherwise recognized by the State to provide these birth attendant services (e.g., doulas, lactation consultant, etc.).*
*For (b) and (c) above, please list and identify below each type of professional who will be providing birth center services: <u>Licensed midwives</u>
TN <u>22-0012</u> Approval Date Effective Date <u>January 1, 2022</u> Supersedes

State/Territory:\_ Louisiana \_

# 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: With limitations** 

II. 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).

#### **Limitations:**

Routine patient costs in qualified clinical trials do not include the following:

- 1. Any investigational item or service that is the subject of the qualifying clinical trial and is not otherwise covered outside of the clinical trial under the State Plan; and
- 2. Any item or service that is provided to the beneficiary solely to satisfy data collection and analysis needs for the qualifying clinical trial and not used in the direct clinical management of the beneficiary and that is not otherwise covered under the State Plan.

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 notconduct or sponsor, and a person is not required to respond to, a collection of information unless it 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>22-0012</u>	_
Supersedes	
TN New Page	

## STATE PLAN UNDER TITLE XIX OF THE SOCIAL SECURITY ACT MEDICAL ASSISTANCE PROGRAM

Attachment 4.19-B Item 30, Page 1

#### STATE OF LOUISIANA

#### PAYMENTS FOR MEDICAL AND REMEDIAL CARE AND SERVICES

METHODS AND STANDARDS FOR ESTABLISHING PAYMENT RATES - OTHER TYPES OF CARE OR SERVICE LISTED IN SECTION 1902(a) OF THE ACT THAT ARE INCLUDED IN THE PROGRAM UNDER THE PLAN ARE DESCRIBED AS FOLLOWS:

1905(a)(30) Qualifying Clinical Trials

#### **Reimbursement Methodology**

Medicaid will reimburse routine patient costs associated with beneficiaries' participating in qualifying clinical trials on or after January 1, 2022.

The following are excluded from reimbursement:

- 1. Any investigational item or service that is the subject of the qualifying clinical trial and is not otherwise covered outside of the clinical trial under the State Plan; and
- 2. Any item or service that is provided to the beneficiary solely to satisfy data collection and analysis needs for the qualifying clinical trial and not used in the direct clinical management of the beneficiary and that is not otherwise covered under the State Plan.

TN <u>22-0012</u>