CONTRACTOR MEDICAL CAMEDIONS OF MICE.	1. TRANSMITTAL NUMBER	2. STATE
TRANSMITTAL AND NOTICE OF APPROVAL OF	1 0 - 0 7	Oklahoma
STATE PLAN MATERIAL FOR: CENTERS FOR MEDICARE & MEDICAID SERVICES	3. PROGRAM IDENTIFICATION: SECURITY ACT (MEDICAID)	
TO: REGIONAL ADMINISTRATOR CENTERS FOR MEDICARE & MEDICAID SERVICES	4. PROPOSED EFFECTIVE DAT	E
DEPARTMENT OF HEALTH AND HUMAN SERVICES	April 1, 2010	
5. TYPE OF PLAN MATERIAL (Check One)		
NEW STATE PLAN AMENDMENT TO BE CONSIDER		X AMENDMENT
COMPLETE BLOCKS 6 THRU 10 IF THIS IS AN AMENDM	7. FEDERAL BUDGET IMPACT	amenoment)
6. FEDERAL STATUTE/REGULATION CITATION	a. FFY 2010	(\$1,061,064) savings
42 CFR 440.120 42 CFR 447.331 42 CFR 447.332	b. FFY 2011	(\$2,138,927) savings
8. PAGE NUMBER OF THE PLAN SECTION OR ATTACHMENT	9. PAGE NUMBER OF THE SUP OR ATTACHMENT (If Applicab	
Attachment 4.19-B Page 7 Attachment 4.19-B Page 7a	Same Page, Revised 10- Same Page, Revised 02-	
10. SUBJECT OF AMENDMENT		
Changes to State Maximum Allowable Cost calculation and  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	X OTHER, AS SPECIFIED The Governor does not material.	
12. SIGNATURE OF STATE AGENCY OFFICIAL	16. RETURN TO	
13. TYPED NAME	Oklahoma Health Care	Authority
Mike Fogarty 14. TITLE	Attn: Cindy Roberts 4545 N. Lincoln Blvd., Suite 124	
Chief Executive Officer	Oklahoma City, OK 73	
15. DATE SUBMITTED	•	
March 25, 2010		
17. DATE RECEIVED  18. D  25. March, 2010  PLAN APPROVED ONE  19. EFFECTIVE DATE OF APPROVED MATERIAL  26.	COPY ATTACKED GENATURE OF REGIONAL OFFICIA	ecember, 2010
21. TYPED NAME BILL Brooks	me Associate Red Div of Med	gional Administrator uchid z Childrens Health
23 REMARKS  C: Mike Fogarty  Girdy Roberts  Tywande Cox  Traylor Rains  Rodney Ikard		

## METHODS AND STANDARDS FOR ESTABLISHING PAYMENT RATES OTHER TYPES OF CARE

#### 6. Payment for prescribed drugs (cont.)

Programming has been developed to review the CMS upper limit products to assure in the aggregate Medicaid expenditures for multiple source drugs do not exceed the federal upper limits. Such reports and all other relevant statistical data are maintained by the Agency and are available on request.

#### **Brand Name Certification**

- (a) When a product is available in both a brand and generic form, a prior authorization is required before the branded product may be dispensed. The prescribing provider must certify the brand name drug product is medically necessary for the well being of the patient, otherwise a generic must be substituted.
  - (1) The certification must be written in the physician's or other prescribing provider's handwriting.
  - (2) It is unacceptable to use a printed box on the prescription blank that could be checked by the physician to indicate brand necessary, or to use a hand-written statement that is transferred to a rubber stamp and then stamped onto the prescription blank.
- (b) The Brand Necessary Certification applies to CMS Federal Upper Limit and State Maximum Allowable Cost (SMAC) products.

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Revised 04-01-10

TN#<u>/0-07</u> Supersedes TN# 02-03 Approval Date 12-22-10

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### State **OKLAHOMA**

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# METHODS AND STANDARDS FOR ESTABLISHING PAYMENT RATES OTHER TYPES OF CARE

6.	Payment f	or prescribed	drugs
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- (a) **Reimbursement**. Reimbursement for pharmacy claims is based on the sum of an estimate of the ingredient cost, plus a dispensing fee.
- (b) Ingredient Cost. Ingredient cost is estimated by one of the following methods:
  - (1) Maximum Allowable Cost.
    - (A) The State Maximum Allowable Cost (SMAC) is established for certain products which have a Food and Drug Administration (FDA) approved generic equivalent. The SMAC is calculated using prices from pharmaceutical wholesalers who supply these products to pharmacy providers in Oklahoma. Pharmacies may challenge a specific product's SMAC price by providing a current invoice that reflects a net cost higher than the calculated SMAC price and by certifying that there is not another product available to them which is generically equivalent to the higher priced product.
    - (B) The Federal Upper Limit (FUL) is established by CMS in accordance with applicable federal laws and regulations.
    - (C) Injectable drugs which are dispensed by a pharmacy through the Vendor Drug Program. Injectable drugs shall be priced based on a formula equivalent to the Medicare Part B allowed charge whether they are furnished through the pharmacy program or through the medical program. Medicare Part B pricing is calculated using Average Sale Price (ASP) plus 6%. When no ASP is available, the Wholesaler Acquisition Cost (WAC) methodology is used. Pricing is updated on a quarterly basis in correspondence with the updates to Medicare Part B pricing.
  - (2) **The Estimated Acquisition Cost**. The Estimated Acquisition Cost (EAC) means the agency's best estimate of the price generally and currently paid by providers for a drug marketed or sold by a particular manufacturer or labeler. EAC is calculated as AWP minus 12%.
- (c) **Maximum allowable dispensing fee.** The maximum allowable dispensing fee for prescribed medication is established by review of surveys. A recommendation is made by the State Plan Amendment Rate Committee and presented to the Oklahoma Health Care Authority Board for their approval. The maximum allowable dispensing fee is \$4.02.
- (d) **Reimbursement for prescription claims.** Prescription claims will be reimbursed using the lower of the following calculation methods:
  - (1) the lower of estimated acquisition cost, Federal Upper Limit (FUL), or State Maximum Allowable Cost (SMAC) plus a dispensing fee, or
  - (2) usual and customary charge to the general public.

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