Authorization Page

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TEMPORARY ADMINISTRATIVE RULES

Oregon Health Authority, Health Systems Division: Medical Assistance Programs	410
Agency and Division	Administrative Rules Chapter Number
Sandy Cafourek	dmap.rules@state.or.us
Rules Coordinator	Email Address
500 Summer St. NE, Salem, OR 97301	503-945-6430
Address	Telephone
Upon filing.	
Adopted on	
10/13/2016 thru 12/27/2016	
Effective dates	

RULE CAPTION

Amending Prior Authorization Approval Criteria Guide

Not more than 15 words

RULEMAKING ACTION

ADOPT:

AMEND: 410-121-0040

SUSPEND:

Stat. Auth.: ORS 413.032, 413.042, 414.065, 414.325, 414.330 to 414.414, 414.312, 414.316

Other Auth.:

Stats. Implemented: ORS 414.065; 414.325, 414.334, 414.361, 414.369, 414.371, 414.353, 414.354

RULE SUMMARY

The Pharmaceutical Services program administrative rules (division 121) govern Division payments for services provided to certain clients. The Authority is amending this rule to update the Oregon Medicaid Fee for Service Prior Authorization Criteria Guide found at http://www.oregon.gov/oha/healthplan/Pages/pharmacy-policy.aspx based on the P&T (Pharmacy and Therapeutic) Committee recommendations.

STATEMENT OF NEED AND JUSTIFICATION

The amendment of OAR 410-121-0040

In the Matter of

414.353, 414.354, and Or Law 2011, chapter 720 (HB 2100),:
http://www.oregon.gov/pers/docs/2011_legislation/hb2100.en.pdf

Documents Relied Upon, and where they are available

The Pharmaceutical Services program administrative rules (division 121) govern Division payments for services provided to certain clients. The Authority needs to amend this rule to update the Oregon Medicaid Fee for Service Prior Authorization Criteria Guide found at http://www.oregon.gov/oha/healthplan/Pages/pharmacy-policy.aspx based on the P&T (Pharmacy and Therapeutic) Committee recommendations.

Need for the Temporary Rule(s)

The Authority finds that failure to act promptly will result in serious prejudice to the public interest, the Authority, and clients enrolled in Oregon's Medicaid Program by delaying the reassessment and update of preferred drug lists and prior authorization requirements. These rules need to be adopted promptly so the Authority can ensure the safe and appropriate use of Medicaid covered drugs.

Justification of Temporary Rules

Authorized Signer	Karen Wheeler	10/13/16
	Printed Name	Date

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Secretary of State

STATEMENT OF NEED AND JUSTIFICATION

A Certificate and Order for Filing Temporary Administrative Rules accompanies this form.

Oregon Health Authority	Health Systems Division (Division)	410
Agency and Division		Administrative Rules Chapter Number

Rule Caption: Amending Prior Authorization Approval Criteria Guide

In the Matter of: The amendment of OAR 410-121-0040

Statutory Authority: ORS 413.032, 413.042, 414.065, 414.325, 414.330 to 414.414, 414.312, 414.316

Other Authority: None

Stats. Implemented: ORS 414.065; 414.325, 414.334, 414.361, 414.369, 414.371, 414.353, 414.354

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[signature]	Karen Wheeler	10/13/16
Authorized Signer	Printed name	Date

Administrative Rules Unit, Archives Division, Secretary of State, 800 Summer Street NE, Salem, Oregon 97310.

410-121-0040

Prior Authorization Required for Drugs and Products

(1) Prescribing practitioners shall obtain prior authorization (PA) for the drugs and categories of drugs requiring PA in this rule, using the procedures set forth in OAR 410-121-0060.

(2) All drugs and categories of drugs including, but not limited to, those drugs and categories of drugs that require PA shall meet the following requirements for coverage:

(a) Each drug shall be prescribed for conditions funded by the Oregon Health Plan (OHP) in a manner consistent with the Health Evidence Review Commission (HERC) Prioritized List of Health Services (OAR 410-141-0480 through 410-141-0520). If the medication is for a non-covered diagnosis, the medication may not be covered unless there is a co-morbid condition for which coverage would be allowed. The use of the medication shall meet corresponding treatment guidelines and be included within the client's benefit package of covered services and not otherwise excluded or limited;

(b) Each drug shall also meet other criteria applicable to the drug or category of drug in these pharmacy provider rules, including PA requirements imposed in this rule.

(3) The Authority may require PA for individual drugs and categories of drugs to ensure that the drugs prescribed are indicated for conditions funded by OHP and consistent with the Prioritized List of Health Services and its corresponding treatment guidelines (see OAR 410-141-0480). The drugs and categories of drugs that the Authority requires PA for this purpose are found in the Oregon Medicaid Fee-for-Service Prior Authorization Approval Criteria (PA Criteria guide) dated October 13~~August 26~~, 2016, adopted and incorporated by reference and found at: http://www.oregon.gov/OHA/healthplan/pages/pharmacy-policy.aspx

(4) The Authority may require PA for individual drugs and categories of drugs to ensure medically appropriate use or to address potential client safety risk associated with the particular drug or category of drug, as recommended by the Pharmacy & Therapeutics Committee (P&T) and adopted by the Authority in this rule. The drugs and categories of drugs for which the Authority requires PA for this purpose are found in the Pharmacy PA Criteria Guide.

(5) New drugs shall be evaluated when added to the weekly upload of the First Databank drug file:

(a) If the new drug is in a class where current PA criteria apply, all associated PA criteria shall be required at the time of the drug file load;

(b) If the new drug is indicated for a condition below the funding line on the Prioritized List of Health Services, PA shall be required to ensure that the drug is prescribed for a condition funded by OHP;

(c) PA criteria for all new drugs shall be reviewed by the DUR/P&T Committee.

(6) PA shall be obtained for brand name drugs that have two or more generically equivalent products available and that are not determined Narrow Therapeutic Index drugs by the DUR/P&T Committee:

(a) Immunosuppressant drugs used in connection with an organ transplant shall be evaluated for narrow therapeutic index within 180 days after United States patent expiration;

(b) Manufacturers of immunosuppressant drugs used in connection with an organ transplant shall notify the Authority of patent expiration within 30 days of patent expiration for section (5)(a) to apply;

(c) Criteria for approval are:

(A) If criteria established in section (3) or (4) of this rule applies, follow that criteria;

(B) If section (6)(A) does not apply, the prescribing practitioner shall document that the use of the generically equivalent drug is medically contraindicated and provide evidence that either the drug has been used and has failed or that its use is contraindicated based on evidence-based peer reviewed literature that is appropriate to the client's medical condition.

(7) PA shall be obtained for non-preferred Preferred Drug List (PDL) products in a class evaluated for the PDL except in the following cases:

(a) The drug is a mental health drug as defined in OAR 410-121-0000;

(b) The original prescription is written prior to 1/1/10;

(c) The prescription is a refill for the treatment of seizures, cancer, HIV, or AIDS; or

(d) The prescription is a refill of an immunosuppressant.

(8) PA may not be required:

(a) When the prescription ingredient cost plus the dispensing fee is less than the PA processing fees as determined by the Authority;

(b) For over-the-counter (OTC) covered drugs when prescribed for conditions covered under OHP; or

(c) If a drug is in a class not evaluated from the Practitioner-Managed Prescription Drug Plan under ORS 414.334.

Stat. Auth.: ORS 413.032, 413.042, 414.065, 414.330 to 414.414, 414.312, 414.316

Stats. Implemented: 414.065, 414.334, 414.361, 414.371, 414.353, 414.354