

Authorization Page

Generated on August 13, 2013 9:40AM

PERMANENT ADMINISTRATIVE RULES

Oregon Health Authority, Division of Medical Assistance  
Programs

410

Agency and Division

Administrative Rules Chapter Number

Cheryl Peters

dmap.rules@state.or.us

Rules Coordinator

Email Address

500 Summer St. NE, Salem, OR 97301

503-945-6527

Address

Telephone

Upon filing.

Adopted on

Upon filing.

Effective date

RULE CAPTION

Amending Preferred Drug List and Prior Authorization Guide-January 31, 2013  
DUR/P&T Action

Not more than 15 words

RULEMAKING ACTION

ADOPT:

AMEND: 410-121-0030, 410-121-0040

REPEAL:

RENUMBER:

AMEND & RENUMBER:

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

Other Auth.: None

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

RULE SUMMARY

The Pharmaceutical Services Program administrative rules (Division 121) govern Division payments for services provided to certain clients. The Division needs to amend rules as follows:

410-121-0030:

- Hyoscyamine rapid tabs preferred
- Oscimin<sup>®</sup> non-preferred
- Hyoscyamine drops non-preferred
- Mirabegron non-preferred
- Gelnique<sup>®</sup> non-preferred
- Tolterodine non-preferred with 90 day grandfather
- Acridinium bromide non-preferred
- Combivent Respimat<sup>®</sup> and Combivent MDI<sup>®</sup> non-preferred with indefinite grandfather
- All erythromycin products non-preferred
- Noroxin<sup>®</sup> non-preferred
- Omeprazole tablets non-preferred
- All OTC H2A products non-preferred
- Cimetidine non-preferred with indefinite grandfather
- All OTC antihistamine products non-preferred

410-121-0040:

Proton Pump Inhibitors update criteria

*Rhonda Busek*

Rhonda Busek

8-15-13

Authorized Signer

Printed Name

Date

Authorization Page replaces the ink signature on paper filings. Have your authorized signer sign and date, then scan and attach it to your filing. You must complete this step before submitting your Permanent and Temporary filings.

## **410-121-0030 Practitioner-Managed Prescription Drug Plan**

(1) The Practitioner-Managed Prescription Drug Plan (PMPDP) is a plan that ensures that fee-for-service clients of the Oregon Health Plan shall have access to the most effective prescription drugs appropriate for their clinical conditions at the best possible price:

(a) Licensed health care practitioners (informed by the latest peer reviewed research), make decisions concerning the clinical effectiveness of the prescription drugs;

(b) The licensed health care practitioners also consider the health condition of a client or characteristics of a client, including the client's gender, race or ethnicity.

(2) PMPDP Preferred Drug List (PDL):

(a) The PDL is the primary tool that the Division developed to inform licensed health care practitioners about the results of the latest peer-reviewed research and cost effectiveness of prescription drugs;

(b) The PDL (as defined in 410-121-0000 (cc) consists of prescription drugs that the Division, in consultation with the Drug Use Review (DUR) / Pharmacy & Therapeutics Committee (P&T), has determined represent the most effective drug(s) available at the best possible price;

(c) The PDL shall include drugs that are Medicaid reimbursable and the Food and Drug Administration (FDA) has determined to be safe and effective.

(3) PMPDP PDL Selection Process:

(a) The Division shall utilize the recommendations made by the P&T, that result from an evidence-based evaluation process, as the basis for selecting the most effective drug(s);

(b) The Division shall determine the drugs selected in (3)(a) that are available for the best possible price and shall consider any input from the P&T about other FDA-approved drug(s) in the same class that are available for a lesser relative price. The Division shall determine relative price using the methodology described in subsection (4);

(c) The Division shall evaluate selected drug(s) for the drug classes periodically:

(A) Evaluation shall occur more frequently at the discretion of the Division if new safety information or the release of new drugs in a class or other information which makes an evaluation advisable;

(B) New drugs in classes already evaluated for the PDL shall be non-preferred until the new drug has been reviewed by the P&T;

(C) The Division shall make all changes or revisions to the PDL, using the rulemaking process and shall publish the changes on the Division's Pharmaceutical Services provider rules Web page.

(4) Relative cost and best possible price determination:

(a) The Division shall determine the relative cost of all drugs in each selected class that are Medicaid reimbursable and that the FDA has determined to be safe and effective;

(b) The Division may also consider dosing issues, patterns of use and compliance issues. The Division shall weigh these factors with any advice provided by the P&T in reaching a final decision;

(5) Pharmacy providers shall dispense prescriptions in the generic form, unless:

(a) The practitioner requests otherwise, subject to the regulations outlined in OAR 410-121-0155;

(b) The brand name medication is listed as preferred on the PDL.

(6) The exception process for obtaining non-preferred physical health drugs that are not on the PDL drugs shall be as follows:

(a) If the prescribing practitioner, in their professional judgment, wishes to prescribe a physical health drug not on the PDL, they may request an exception, subject to the requirements of OAR 410-121-0040;

(b) The prescribing practitioner must request an exception for physical health drugs not listed in the PDL subject to the requirements of OAR 410-121-0060;

(c) Exceptions shall be granted in instances:

(A) Where the prescriber in their professional judgment determines the non-preferred drug is medically appropriate after consulting with the Division or the Oregon Pharmacy Help Desk; or

(B) Where the prescriber requests an exception subject to the requirement of (6)(b) and fails to receive a report of PA status within 24 hours, subject to OAR 410-121-0060.

(7) Table 121-0030-1, PMPDP PDL dated May 1, 2013 is incorporated in rule by reference and is found on our Web page at [www.orpdl.org](http://www.orpdl.org).

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

Stats. Implemented: ORS 414.065, 414.325, 414.334, 414.361, 414.369 & 414.371

#### **410-121-0040 Prior Authorization Required for Drugs and Products**

(1) Prescribing practitioners are responsible for obtaining prior authorization (PA) for the drugs and categories of drugs requiring PA in this rule, using the procedures required 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 as described in this rule, are subject to the following requirements for coverage:

(a) Each drug must be prescribed for conditions funded by Oregon Health Plan (OHP) in a manner consistent with the Oregon Health Services Commission's Prioritized List of Health Services (OAR 410141-0480 through 410-141-0520). If the medication is for a non-covered diagnosis, the medication shall not be covered unless there is a co-morbid condition for which coverage would be extended. The use of the medication must meet corresponding treatment guidelines, be included within the client's benefit package of covered services, and not otherwise excluded or limited;

(b) Each drug must 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 Oregon Health Authority (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 OHP Fee-For-Service Pharmacy PA Criteria Guide (PA Criteria Guide) dated May 1, 2013, incorporated in rule by reference and found on our Web page at:  
<http://www.dhs.state.or.us/policy/healthplan/guides/pharmacy/clinical.html>

(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 (see OAR 410-121-0100 for a description of the DUR program). 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 is required for brand name drugs that have two or more generically equivalent products available and that are NOT determined Narrow Therapeutic Index drugs by the Oregon DUR/P&T Committee:

(a) Immunosuppressant drugs used in connection with an organ transplant must 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 must notify the department of patent expiration within 30 days of patent expiration for (5)(a) to apply;

(c) Criteria for approval are:

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

(B) If (6)(A) does not apply, the prescribing practitioner must 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 is required 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, 414.325, & 414.330 to 414.414

Stats. Implemented: 414.065, 414.325, 414.334, 414.361, 414.369 & 414.371