



121st MAINE LEGISLATURE

FIRST REGULAR SESSION-2003

Legislative Document

No. 1288

S.P. 419

In Senate, March 11, 2003

An Act To Increase Public Access to the Prior Authorization Process

Reference to the Committee on Health and Human Services suggested and ordered printed.

JOY J. O'BRIEN Secretary of the Senate

Presented by Senator TURNER of Cumberland. Cosponsored by Representative DUGAY of Cherryfield and Senators: BRENNAN of Cumberland, MARTIN of Aroostook, WESTON of Waldo, Representatives: CURLEY of Scarborough, KANE of Saco.

	Be it enacted by the People of the State of Maine as follows:
2	Sec. 1. 22 MRSA §3174-DD is enacted to read:
4	<u>\$3174-DD. Drug Utilization Review Committee</u>
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8	1. Authority. The department has the authority to determine which prescription and over-the-counter drugs are subject to prior authorization and coverage under the MaineCare
10	program.
12	2. Drug Utilization Review Committee. In order to make determinations regarding which prescription and over-the-counter
14	drugs are subject to prior authorization, the Drug Utilization Review Committee, referred to in this section as "the committee,"
16	is established.
18	A. The committee is composed of an equal number of actively practicing physicians and actively practicing pharmacists,
20	totaling not fewer than 6 individuals. The department shall appoint the members of the committee and establish their
22	terms.
24	B. Public notice of the date, time and location of all meetings of the committee must be made in accordance with
26	Title 1, section 406 at least 7 days in advance of the meeting. This notice must also include a list of all drugs
28	to be considered for prior authorization at that meeting. The meetings must be considered public proceedings, for
30	purposes of Title 1, chapter 13. Members of the public, including health care providers and drug manufacturers, who
32	are physically present at committee meetings must be granted reasonable opportunity to address the committee prior to any
34	vote.
36	C. At the conclusion of a meeting pursuant to paragraph B, committee members shall record their vote for each drug on a
38	form to be supplied at each meeting. After adding or deleting a drug from the list of drugs that require prior
40	authorization, the committee shall issue written findings setting forth the evidentiary basis for its decision. These
42	findings must address, without limitation, all relevant clinical data, as well as the likelihood that subjecting a
44	<u>drug to prior authorization will cause adverse medical</u> results or affect patients' access to all medically
46	necessary outpatient drugs. A vote of at least 2/3 of the committee members present is required to add or delete a
48	drug from the list of drugs that require prior authorization.

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D. All documents generated by the committee are public records for purposes of Title 1, chapter 13 and must be maintained on file at the Bureau of Medical Services. 3. Appeals. A determination that a particular drug is

 6 subject to prior authorization constitutes final agency action within the meaning of the Maine Administrative Procedure Act and
8 is subject to judicial review under the Maine Administrative Procedure Act. Persons entitled to appeal a decision of the
10 committee to subject a particular drug to prior authorization are limited to:

A. MaineCare recipients who were being prescribed the drug at the time of the committee's decision;

16 <u>B. Health care providers who had prescribed the drug at the time of the committee's decision; and</u>

C. The manufacturer of the drug.

 4. Rules. The department shall adopt rules to implement
22 this section. Rules adopted pursuant to this subsection are routine technical rules as defined in Title 5, chapter 375,
24 subchapter 2-A.

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SUMMARY

bill This establishes the Drug Utilization Review 30 Committee. The committee makes determinations regarding which prescription and over-the-counter drugs are subject to prior 32 authorization under the MaineCare program. The bill requires public notice of committee meetings to be given, and provides that committee meetings are public proceedings and committee 34 documents are public records for purposes of the laws governing freedom of access. It provides that members of the public must 36 be granted a reasonable opportunity to address the committee and requires the committee to issue written findings that describe 38 the basis for its decisions. It also provides that a 2/3 vote of 40 the committee is required to add or delete a drug from the list of drugs that require prior authorization and that the decisions 42 of the committee are final agency action for purposes of the Maine Administrative Procedure Act.