## First Regular Session Sixty-ninth General Assembly STATE OF COLORADO

## PREAMENDED

This Unofficial Version Includes Committee Amendments Not Yet Adopted on Second Reading

LLS NO. 13-0886.01 Christy Chase x2008

SENATE BILL 13-277

#### SENATE SPONSORSHIP

Aguilar, Morse

### **HOUSE SPONSORSHIP**

**Ginal,** Duran, Fields, Garcia, Gerou, Hamner, Hullinghorst, Joshi, Melton, Moreno, Peniston, Pettersen, Primavera, Rosenthal, Ryden, Salazar, Schafer, Singer, Williams

Senate Committees Health & Human Services House Committees Health, Insurance & Environment Appropriations

# A BILL FOR AN ACT

101	CONCERNING THE DEVELOPMENT OF A PRIOR AUTHORIZATION
102	PROCESS TO BE USED IN OBTAINING PRIOR APPROVAL FROM
103	CARRIERS FOR COVERAGE OF DRUG BENEFITS, AND, IN
104	CONNECTION THEREWITH, MAKING AN APPROPRIATION.

#### **Bill Summary**

(Note: This summary applies to this bill as introduced and does not reflect any amendments that may be subsequently adopted. If this bill passes third reading in the house of introduction, a bill summary that applies to the reengrossed version of this bill will be available at http://www.leg.state.co.us/billsummaries.)

The bill requires the commissioner of insurance (commissioner) to develop, by July 31, 2014, and prescribing providers, carriers, and, if



SENATE Amended 2nd Reading April 29, 2013 applicable, pharmacy benefit management firms (PBMs), to use, by January 1, 2015, a uniform prior authorization process for purposes of submitting and receiving requests for prior coverage approval of a drug benefit.

The commissioner is directed to adopt rules to establish the prior authorization process, which is to include specified components aimed at creating uniformity and reducing administrative burdens on prescribing providers, carriers, and PBMs, as well as making the criteria used for deciding prior authorization requests transparent and establishing a procedure for waiving the process under extenuating circumstances.

To assist in developing the process, the commissioner is to appoint a work group of various stakeholders to make recommendations on specified aspects of the process that the commissioner is to consider, including national standards for electronic prior authorization.

Once the prior authorization process is established, the request is deemed granted if a carrier or PBM fails to use or accept the prior authorization process, fails to notify the prescribing provider within a specified period that the request is approved or denied or that additional information is required to process the request, or fails to notify the prescribing provider within a specified period after receipt of the required additional information that the request is approved or denied. An approved prior authorization is valid for at least 180 days after the date of approval.

1 Be it enacted by the General Assembly of the State of Colorado:

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#### **SECTION 1. Legislative declaration.** (1) The general assembly

3 hereby finds that:

4 (a) Carriers and pharmacy benefit management firms routinely
5 require health care providers to request prior authorization when
6 prescribing medications or treatments not routinely covered by health
7 plan formularies;

8 (b) Each carrier and pharmacy benefit management firm has its 9 own prior authorization process, and the multiplicity of prior 10 authorization processes imposes a significant administrative burden on 11 health care providers, resulting in delayed patient access to medication 12 and increased administrative costs; and (c) A standardized prior authorization process that any health care
 provider can use, regardless of the carrier, pharmacy benefit management
 firm, or health plan that covers that provider's patient, will simplify the
 administrative process and improve patient care by allowing health care
 providers to devote less time to administrative duties and more time to
 patient care.

7 SECTION 2. In Colorado Revised Statutes, add 10-16-124.5 as
8 follows:

9 **10-16-124.5.** Prior authorization form - drug benefits - rules 10 of commissioner - definition. (1) (a) NOTWITHSTANDING ANY OTHER 11 PROVISION OF LAW BUT SUBJECT TO PARAGRAPH (b) OF THIS SUBSECTION 12 (1), ON AND AFTER JANUARY 1, 2015, A CARRIER OR, IF A CARRIER 13 CONTRACTS WITH A PHARMACY BENEFIT MANAGEMENT FIRM TO PERFORM 14 PRIOR AUTHORIZATION SERVICES FOR DRUG BENEFITS, THE PHARMACY 15 BENEFIT MANAGEMENT FIRM, SHALL UTILIZE THE PRIOR AUTHORIZATION 16 PROCESS DEVELOPED PURSUANT TO SUBSECTION (3) OF THIS SECTION 17 WHEN REQUIRING PRIOR AUTHORIZATION FOR DRUG BENEFITS.

(b) This section does not apply to a nonprofit health
MAINTENANCE ORGANIZATION WITH RESPECT TO MANAGED CARE PLANS
THAT PROVIDE A MAJORITY OF COVERED PROFESSIONAL SERVICES
THROUGH A SINGLE CONTRACTED MEDICAL GROUP.

(2) (a) EXCEPT AS PROVIDED IN PARAGRAPH (b) OF THIS
 SUBSECTION (2), A PRIOR AUTHORIZATION REQUEST IS DEEMED GRANTED
 IF A CARRIER OR PHARMACY BENEFIT MANAGEMENT FIRM FAILS TO:

25 (I) UTILIZE THE PRIOR AUTHORIZATION PROCESS DEVELOPED
 26 PURSUANT TO SUBSECTION (3) OF THIS SECTION;

27 (II) FOR PRIOR AUTHORIZATION REQUESTS SUBMITTED

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1 ELECTRONICALLY:

(A) NOTIFY THE PRESCRIBING PROVIDER, WITHIN TWO BUSINESS
DAYS AFTER RECEIPT OF THE REQUEST, THAT THE REQUEST IS APPROVED,
DENIED, OR INCOMPLETE, AND IF INCOMPLETE, INDICATE THE SPECIFIC
ADDITIONAL INFORMATION, CONSISTENT WITH CRITERIA POSTED
PURSUANT TO SUBPARAGRAPH (II) OF PARAGRAPH (a) OF SUBSECTION (3)
OF THIS SECTION, THAT IS REQUIRED TO PROCESS THE REQUEST; OR

8 (B) NOTIFY THE PRESCRIBING PROVIDER, WITHIN TWO BUSINESS 9 DAYS AFTER RECEIVING THE ADDITIONAL INFORMATION REQUIRED BY THE 10 CARRIER OR PHARMACY BENEFIT MANAGEMENT FIRM PURSUANT TO 11 <u>SUB-SUBPARAGRAPH (A) OF THIS SUBPARAGRAPH (II),</u> THAT THE REQUEST 12 IS APPROVED OR DENIED;

13 (III) FOR NONURGENT PRIOR AUTHORIZATION REQUESTS
14 SUBMITTED ORALLY OR BY FACSIMILE OR ELECTRONIC MAIL, NOTIFY THE
15 PRESCRIBING PROVIDER, WITHIN <u>THREE BUSINESS DAYS</u> AFTER RECEIPT OF
16 THE REQUEST, THAT THE REQUEST IS APPROVED OR DENIED; AND

17 (IV) FOR URGENT PRIOR AUTHORIZATION REQUESTS SUBMITTED
18 ORALLY OR BY FACSIMILE OR ELECTRONIC MAIL, NOTIFY THE PRESCRIBING
19 PROVIDER, WITHIN <u>ONE DAY</u> AFTER RECEIPT OF THE REQUEST, THAT THE
20 REQUEST IS APPROVED OR DENIED.

(b) IF A CARRIER OR PHARMACY BENEFIT MANAGEMENT FIRM
 NOTIFIES THE PRESCRIBING PROVIDER PURSUANT TO SUB-SUBPARAGRAPH
 (A) OF SUBPARAGRAPH (II) OF PARAGRAPH (a) OF THIS SUBSECTION (2)
 THAT A PRIOR AUTHORIZATION REQUEST IS INCOMPLETE AND THAT
 ADDITIONAL INFORMATION IS REQUIRED, THE PRESCRIBING PROVIDER
 SHALL SUBMIT THE ADDITIONAL INFORMATION WITHIN TWO BUSINESS
 DAYS AFTER RECEIPT OF THE NOTICE FROM THE CARRIER OR PHARMACY

1 BENEFIT MANAGEMENT FIRM. IF THE PRESCRIBING PROVIDER FAILS TO 2 SUBMIT THE REQUIRED ADDITIONAL INFORMATION WITHIN TWO BUSINESS 3 DAYS AFTER RECEIPT OF THE NOTICE, THE REQUEST IS NOT DEEMED 4 GRANTED PURSUANT TO PARAGRAPH (a) OF THIS SUBSECTION (2). AFTER 5 RECEIPT OF THE REQUIRED ADDITIONAL INFORMATION, THE CARRIER OR 6 PHARMACY BENEFIT MANAGEMENT FIRM SHALL RESPOND TO THE PRIOR 7 AUTHORIZATION REQUEST IN ACCORDANCE WITH SUB-SUBPARAGRAPH (B)8 OF SUBPARAGRAPH (II) OF PARAGRAPH (a) OF THIS SUBSECTION (2). 9 (3) (a) ON OR BEFORE JULY 31, 2014, THE COMMISSIONER SHALL 10 DEVELOP, BY RULE, A UNIFORM PRIOR AUTHORIZATION PROCESS THAT:

(I) IS MADE AVAILABLE ELECTRONICALLY BY THE CARRIER OR
 PHARMACY BENEFIT MANAGEMENT FIRM BUT THAT DOES NOT REQUIRE THE
 PRESCRIBING PROVIDER TO SUBMIT A PRIOR AUTHORIZATION REQUEST
 ELECTRONICALLY;

(II) REQUIRES EACH CARRIER AND PHARMACY BENEFIT
MANAGEMENT FIRM TO MAKE THE FOLLOWING AVAILABLE AND
ACCESSIBLE IN A CENTRALIZED LOCATION ON ITS WEB SITE:

18 (A) ITS PRIOR AUTHORIZATION REQUIREMENTS AND RESTRICTIONS,
19 INCLUDING A LIST OF DRUGS THAT REQUIRE PRIOR AUTHORIZATION;

(B) WRITTEN CLINICAL CRITERIA THAT ARE EASILY
UNDERSTANDABLE TO THE PRESCRIBING PROVIDER AND THAT INCLUDE THE
CLINICAL CRITERIA FOR REAUTHORIZATION OF A PREVIOUSLY APPROVED
DRUG AFTER THE PRIOR AUTHORIZATION PERIOD HAS EXPIRED; AND

24 (C) THE STANDARD FORM FOR SUBMITTING REQUESTS;

(III) ENSURES THAT CARRIERS AND PHARMACY BENEFIT
MANAGEMENT FIRMS USE EVIDENCE-BASED GUIDELINES, WHEN POSSIBLE,
WHEN MAKING PRIOR AUTHORIZATION DETERMINATIONS;

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(IV) PERMITS, BUT DOES NOT REQUIRE, A PRESCRIBING PROVIDER
 TO SUBMIT A REQUEST FOR A PRIOR AUTHORIZATION FOR DRUG BENEFITS
 ELECTRONICALLY TO THE CARRIER OR PHARMACY BENEFIT MANAGEMENT
 FIRM;

5 (V) REQUIRES CARRIERS AND PHARMACY BENEFIT MANAGEMENT 6 FIRMS, WHEN NOTIFYING THE PRESCRIBING PROVIDER OF ITS DECISION TO 7 APPROVE A PRIOR AUTHORIZATION REQUEST, TO INCLUDE IN THE NOTICE 8 A UNIQUE PRIOR AUTHORIZATION NUMBER ATTRIBUTABLE TO THE 9 PARTICULAR REQUEST, SPECIFICATION OF THE PARTICULAR DRUG BENEFIT 10 APPROVED, THE NEXT DATE FOR REVIEW OF THE APPROVED DRUG BENEFIT, 11 AND A LINK TO THE CURRENT CRITERIA THAT THE PRESCRIBING PROVIDER 12 WILL NEED TO SUBMIT FOR REAPPROVAL OF THE PRIOR AUTHORIZATION; 13 AND

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15 (VI) REQUIRES CARRIERS AND PHARMACY BENEFIT MANAGEMENT
16 FIRMS, WHEN NOTIFYING A PRESCRIBING PROVIDER OF ITS DECISION TO
17 DENY A PRIOR AUTHORIZATION REQUEST, TO INCLUDE A NOTICE THAT THE
18 COVERED PERSON HAS A RIGHT TO APPEAL THE ADVERSE DETERMINATION
19 PURSUANT TO SECTIONS 10-16-113 AND <u>10-16-113.5.</u>

(b) IN DEVELOPING THE UNIFORM PRIOR AUTHORIZATION PROCESS,
THE COMMISSIONER SHALL TAKE INTO CONSIDERATION THE
RECOMMENDATIONS, IF ANY, OF THE WORK GROUP ESTABLISHED
PURSUANT TO SUBSECTION (4) OF THIS SECTION AND THE FOLLOWING:

25 (I) NATIONAL STANDARDS PERTAINING TO ELECTRONIC PRIOR
 26 <u>AUTHORIZATION, INCLUDING, BUT NOT LIMITED TO, STANDARDS</u>
 27 REFERENCED IN FEDERAL LAW;

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(II) WHETHER THE PRIOR AUTHORIZATION PROCESS SHOULD
 REQUIRE CARRIERS AND PHARMACY BENEFIT MANAGEMENT FIRMS, WHEN
 REVIEWING A PRIOR AUTHORIZATION REQUEST, TO USE CLEARLY
 ACCESSIBLE, CONSISTENTLY APPLIED, AND WRITTEN CLINICAL CRITERIA
 BASED ON MEDICAL NECESSITY OR THE APPROPRIATENESS OF THE DRUG
 BENEFIT FOR THE COVERED PERSON;

7 (III) WHETHER THE PRIOR AUTHORIZATION PROCESS SHOULD
8 REQUIRE CARRIERS TO TAKE INTO ACCOUNT, IN DETERMINING CRITERIA
9 FOR PRIOR AUTHORIZATIONS, THE COLORADO PART B MEDICARE
10 CONTRACTOR LOCAL COVERAGE DETERMINATIONS, THE FEDERAL CENTERS
11 FOR MEDICARE AND MEDICAID SERVICES NATIONAL COVERAGE
12 DETERMINATIONS, AND SPECIALTY SOCIETY GUIDELINES, SUCH AS THOSE
13 OF THE AMERICAN SOCIETY OF CLINICAL ONCOLOGY; AND

(IV) WHETHER CARRIERS AND PHARMACY BENEFIT MANAGEMENT
FIRMS COULD USE A RULES ENGINE WITH CRITERIA-DRIVEN QUESTIONS
THAT LEAD TO AN IMMEDIATE DETERMINATION OF A PRIOR
AUTHORIZATION REQUEST OR REQUEST FOR SUBMITTAL OF SPECIFIC
ADDITIONAL INFORMATION NEEDED TO MAKE THE DETERMINATION.

19 (c) IN ADDITION TO THE PRIOR AUTHORIZATION PROCESS, THE 20 COMMISSIONER SHALL DEVELOP, BY RULE, A STANDARDIZED PRIOR 21 AUTHORIZATION FORM, NOT TO EXCEED TWO PAGES IN LENGTH, FOR USE 22 IN SUBMITTING ELECTRONIC AND NONELECTRONIC PRIOR AUTHORIZATION 23 REQUESTS. IN DEVELOPING THE FORM, THE COMMISSIONER SHALL TAKE 24 INTO CONSIDERATION EXISTING FORMS, INCLUDING EXISTING PRIOR 25 AUTHORIZATION FORMS ESTABLISHED BY THE FEDERAL CENTERS FOR 26 MEDICARE AND MEDICAID SERVICES OR THE DEPARTMENT OF HEALTH 27 CARE POLICY AND FINANCING.

(4) (a) WITHIN THIRTY DAYS AFTER THE EFFECTIVE DATE OF THIS
 SECTION, THE COMMISSIONER SHALL ESTABLISH A WORK GROUP
 COMPRISED OF REPRESENTATIVES OF:

4 (I) THE DEPARTMENT OF REGULATORY AGENCIES;

5 (II) LOCAL AND NATIONAL CARRIERS;

6 (III) CAPTIVE AND NONCAPTIVE PHARMACY BENEFIT
7 MANAGEMENT FIRMS;

8 (IV) PROVIDERS, INCLUDING HOSPITALS, PHYSICIANS, ADVANCED
9 PRACTICE NURSES WITH PRESCRIPTIVE AUTHORITY, AND PHARMACISTS;

10 (V) DRUG MANUFACTURERS;

11 (VI) MEDICAL PRACTICE MANAGERS;

12 (VII) CONSUMERS; AND

13 (VIII) OTHER STAKEHOLDERS DEEMED APPROPRIATE BY THE
14 COMMISSIONER.

15 (b) THE WORK GROUP SHALL ASSIST THE COMMISSIONER IN 16 DEVELOPING THE PRIOR AUTHORIZATION PROCESS AND SHALL MAKE 17 RECOMMENDATIONS TO THE COMMISSIONER ON THE ITEMS SET FORTH IN 18 PARAGRAPH (b) OF SUBSECTION (3) OF THIS SECTION. THE WORK GROUP 19 SHALL REPORT ITS RECOMMENDATIONS TO THE COMMISSIONER NO LATER 20 THAN SIX MONTHS AFTER THE COMMISSIONER APPOINTS THE WORK GROUP 21 MEMBERS. REGARDLESS OF WHETHER THE WORK GROUP SUBMITS 22 RECOMMENDATIONS TO THE COMMISSIONER, THE COMMISSIONER SHALL 23 NOT DELAY OR EXTEND THE DEADLINE FOR THE ADOPTION OF RULES 24 CREATING THE PRIOR AUTHORIZATION PROCESS AS SPECIFIED IN 25 PARAGRAPH (a) OF SUBSECTION (3) OF THIS SECTION.

26 (5) NOTWITHSTANDING ANY OTHER PROVISION OF LAW, <u>ON AND</u>
 27 <u>AFTER</u> JANUARY 1, 2015, EVERY PRESCRIBING PROVIDER SHALL USE THE

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PRIOR AUTHORIZATION PROCESS DEVELOPED PURSUANT TO SUBSECTION
 (3) OF THIS SECTION TO REQUEST PRIOR AUTHORIZATION FOR COVERAGE
 OF DRUG BENEFITS, AND EVERY CARRIER AND PHARMACY BENEFIT
 MANAGEMENT FIRM SHALL USE THAT PROCESS FOR PRIOR AUTHORIZATION
 FOR DRUG BENEFITS.

6 (6) UPON APPROVAL BY THE CARRIER OR PHARMACY BENEFIT 7 MANAGEMENT FIRM, A PRIOR AUTHORIZATION IS VALID FOR AT LEAST ONE 8 HUNDRED EIGHTY DAYS AFTER THE DATE OF APPROVAL. IF, AS A RESULT 9 OF A CHANGE TO THE CARRIER'S FORMULARY, THE DRUG FOR WHICH THE 10 CARRIER OR PHARMACY BENEFIT MANAGEMENT FIRM HAS PROVIDED PRIOR 11 AUTHORIZATION IS REMOVED FROM THE FORMULARY OR MOVED TO A LESS 12 PREFERRED TIER STATUS, THE CHANGE IN THE STATUS OF THE PREVIOUSLY 13 APPROVED DRUG DOES NOT AFFECT A COVERED PERSON WHO RECEIVED 14 PRIOR AUTHORIZATION BEFORE THE EFFECTIVE DATE OF THE CHANGE FOR 15 THE REMAINDER OF THE COVERED PERSON'S PLAN YEAR. NOTHING IN THIS 16 SUBSECTION (6) LIMITS THAT ABILITY OF A CARRIER OR PHARMACY 17 BENEFIT MANAGEMENT FIRM, IN ACCORDANCE WITH THE TERMS OF THE 18 HEALTH BENEFIT PLAN, TO SUBSTITUTE A GENERIC DRUG, WITH THE 19 PRESCRIBING PROVIDER'S APPROVAL AND PATIENT'S CONSENT, FOR A 20 PREVIOUSLY APPROVED BRAND-NAME DRUG.

(7) FOR PURPOSES OF THIS SECTION, A PRIOR AUTHORIZATION
REQUEST IS SUBMITTED "ELECTRONICALLY" IF THE PRESCRIBING PROVIDER
SUBMITS THE REQUEST TO THE CARRIER OR PHARMACY BENEFIT
MANAGEMENT FIRM THROUGH A SECURE, WEB-BASED INTERNET PORTAL.
A PRIOR AUTHORIZATION REQUEST SUBMITTED BY ELECTRONIC MAIL IS
NOT SUBMITTED "ELECTRONICALLY".

27 (8) AS USED IN THIS SECTION:

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1 (a) "PRESCRIBING PROVIDER" MEANS A PROVIDER WHO IS: 2 (I) AUTHORIZED BY LAW TO PRESCRIBE ANY DRUG OR DEVICE TO 3 TREAT A MEDICAL CONDITION OF A COVERED PERSON; AND 4 (II) ACTING WITHIN THE SCOPE OF THAT AUTHORITY. 5 (b) "URGENT PRIOR AUTHORIZATION REQUEST" MEANS A REQUEST 6 FOR PRIOR AUTHORIZATION OF A DRUG BENEFIT THAT, BASED ON THE 7 REASONABLE OPINION OF THE PRESCRIBING PROVIDER WITH KNOWLEDGE 8 OF THE COVERED PERSON'S MEDICAL CONDITION, IF DETERMINED IN THE 9 TIME ALLOWED FOR NONURGENT PRIOR AUTHORIZATION REQUESTS, 10 COULD: 11 (I) SERIOUSLY JEOPARDIZE THE LIFE OR HEALTH OF THE COVERED 12 PERSON OR THE ABILITY OF THE COVERED PERSON TO REGAIN MAXIMUM 13 FUNCTION: OR 14 (II) SUBJECT THE COVERED PERSON TO SEVERE PAIN THAT CANNOT 15 BE ADEQUATELY MANAGED WITHOUT THE DRUG BENEFIT THAT IS THE 16 SUBJECT OF THE PRIOR AUTHORIZATION REQUEST. 17 SECTION 3. Appropriation. In addition to any other 18 appropriation, there is hereby appropriated, out of any moneys in the 19 division of insurance cash fund created in section 10-1-103 (3), Colorado 20 Revised Statutes, not otherwise appropriated, to the department of 21 regulatory agencies, for the fiscal year beginning July 1, 2013, the sum of 22 \$8,756 and 0.1 FTE, or so much thereof as may be necessary, for 23 allocation to the division of insurance for personal services related to the 24 implementation of this act. 25

25 SECTION <u>4.</u> Safety clause. The general assembly hereby finds,
 26 determines, and declares that this act is necessary for the immediate
 27 preservation of the public peace, health, and safety.