STATE OF NEW YORK

463

2023-2024 Regular Sessions

IN ASSEMBLY

January 9, 2023

Introduced by M. of A. McDONALD, STECK -- read once and referred to the Committee on Insurance

AN ACT to amend the insurance law and the public health law, in relation to requiring notice of adverse step therapy determinations

The People of the State of New York, represented in Senate and Assembly, do enact as follows:

1	Section 1. Subsection (a) of section 4902 of the insurance law is
2	amended by adding a new paragraph 14 to read as follows:
3	(14) Establishment of a written procedure to assure that the notice of
4	an adverse determination in relation to a step therapy protocol override
5	determination request includes:
б	(i) the reasons for the determination including the clinical ration-
7	<u>ale, if any;</u>
8	(ii) instructions on how to initiate standard and expedited appeals
9	pursuant to section four thousand nine hundred four of this article and
10	an external appeal pursuant to section four thousand nine hundred four-
11	teen of this article;
12	(iii) information that includes: any applicable alternative covered
13	medications; the clinical review criteria relied upon to make such
14	determination; and any additional necessary information that must be
15	provided to, or obtained by, the utilization review agent in order to
16	render a decision on the appeal.
17	§ 2. Section 4903 of the insurance law is amended by adding a new
18	subsection (e-1) to read as follows:
19	(e-1) Notice of an adverse determination made by a utilization review
20	agent in relation to a step therapy protocol override determination
21	request shall be made in writing to the insured or the insured's author-
22	ized representative and the insured's prescribing health care profes-
23	sional as defined in subsection (f) of section forty-nine hundred of
24	this chapter, and shall include:

EXPLANATION--Matter in <u>italics</u> (underscored) is new; matter in brackets [-] is old law to be omitted.

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A. 463

1	(i) the reasons for the determination including the clinical ration-
2	ale, if any;
3	(ii) instructions on how to initiate standard and expedited appeals
4	pursuant to section four thousand nine hundred four of this article and
5	an external appeal pursuant to section four thousand nine hundred four-
б	teen of this article;
7	(iii) information that includes: any applicable alternative covered
8	medications; the clinical review criteria relied upon to make such
9	determination; and any additional necessary information that must be
10	provided to, or obtained by, the utilization review agent in order to
11	render a decision on the appeal.
12	§ 3. Subdivision 1 of section 4902 of the public health law is amended
13	by adding a new paragraph (1) to read as follows:
14	(1) Establishment of a written procedure to assure that the notice of
15	an adverse determination in relation to a step therapy protocol override
16	determination request includes: (i) the reasons for the determination
17	including the clinical rationale, if any; (ii) instructions on how to
18	initiate standard and expedited appeals pursuant to subdivision two of
19	section forty-nine hundred four of this title and an external appeal
20	pursuant to section forty-nine hundred fourteen of this article; (iii)
21	information that includes: any applicable alternative covered medica-
22	tions; the clinical review criteria relied upon to make such determi-
23	nation; and any additional necessary information that must be provided
24	to, or obtained by, the utilization review agent in order to render a
25	decision on the appeal.
26	§ 4. Section 4903 of the public health law is amended by adding a new
27	subdivision 5-a to read as follows:
28	5-a. Notice of an adverse determination made by a utilization review
29	agent in relation to a step therapy protocol override determination
30	request shall be made in writing to the enrollee or the enrollee's
31	authorized representative and the enrollee's prescribing health care
32	professional as defined in subdivision six of section forty-nine hundred
33	of this title, and shall include:
34	(a) the reasons for the determination including the clinical ration-
35	<u>ale, if any;</u>
36	(b) instructions on how to initiate standard and expedited appeals
37	pursuant to section forty-nine hundred four of this title and an
38	external appeal pursuant to section forty-nine hundred fourteen of this
39	<u>article;</u>
40	(c) information that includes: any applicable alternative covered
41	medications; the clinical review criteria relied upon to make such
42	determination; and any additional necessary information that must be
43	provided to, or obtained by, the utilization review agent in order to
44	render a decision on the appeal.
45	§ 5. This act shall take effect on the ninetieth day after it shall
46	have become a law. Effective immediately, the addition, amendment and/or
47	repeal of any rule or regulation necessary for the implementation of
48	this act on its effective date are authorized to be made and completed
49	on or before such effective date.