
THE GENERAL ASSEMBLY OF PENNSYLVANIA

SENATE BILL

No. 1187 Session of
2018

INTRODUCED BY WHITE, SCARNATI, WARD, LANGERHOLC, REGAN, FOLMER,
BARTOLOTTA, VULAKOVICH, RESCHENTHALER, BROWNE, ARGALL,
KILLION, YAW, AUMENT AND STEFANO, JUNE 1, 2018

REFERRED TO LABOR AND INDUSTRY, JUNE 1, 2018

AN ACT

1 Amending the act of June 2, 1915 (P.L.736, No.338), entitled "An
2 act defining the liability of an employer to pay damages for
3 injuries received by an employe in the course of employment;
4 establishing an elective schedule of compensation; providing
5 procedure for the determination of liability and compensation
6 thereunder; and prescribing penalties," in liability and
7 compensation, further providing for prescription drugs and
8 the treatment of work-related injuries; and, in procedure,
9 further providing for peer review.

10 The General Assembly of the Commonwealth of Pennsylvania
11 hereby enacts as follows:

12 Section 1. Section 306(f.1) (3) (vi) (A) and (6) (ii) of the act
13 of June 2, 1915 (P.L.736, No.338), known as the Workers'
14 Compensation Act, are amended, paragraph (3) (vi) is amended by
15 adding subclauses and paragraph (6) is amended by adding a
16 subparagraph to read:

17 Section 306. The following schedule of compensation is
18 hereby established:

19 * * *

20 (f.1) * * *

21 (3) * * *

1 (vi) (A) [The] Except as otherwise provided in subclause
2 (A.1), the reimbursement for prescription drugs and professional
3 pharmaceutical services shall be limited to one hundred ten per
4 centum of the average wholesale price (AWP) of the product,
5 calculated on a per unit basis, as of the date of dispensing.

6 (A.1) All prescriptions of compounded drugs, reimbursement
7 shall be limited to one hundred ten per centum of the average
8 wholesale price of each ingredient in the compounded drug, based
9 on the original manufacturers National Drug Code (NDC) number.

10 * * *

11 (J) The department, in consultation with the Department of
12 Health and the Department of Drug and Alcohol Programs and the
13 appropriate licensing boards under the Department of State,
14 shall develop evidence-based prescription guidelines appropriate
15 to the prescription of pain medication drugs prescribed for or
16 related to the treatment of work-related injuries, including,
17 but not limited to, the type, dosage and duration of
18 prescriptions, and specifically including opioid medications and
19 any other pain medications the departments recognize as
20 potentially addictive. The following shall apply:

21 (I) Within thirty (30) days of the effective date of this
22 subclause, the department shall solicit public comments
23 regarding the development of these prescription guidelines. The
24 public comment period shall be ninety (90) days, during which
25 time the department shall also hold at least three public
26 hearings. The department shall publish notice of the public
27 comment period and public hearings in the Pennsylvania Bulletin.

28 (II) Within thirty (30) days after the close of the public
29 comment period under subprovision (I), the department shall
30 publish these prescription guidelines in the Pennsylvania

1 Bulletin. The prescription guidelines shall apply to any
2 prescription of pain medication drugs for or related to the
3 treatment of work-related injuries made after thirty (30) days
4 from the date of the publication.

5 (III) In developing the evidence-based prescription
6 guidelines, the department shall consider the following factors:

7 (a) That the guidelines assure injured workers access to
8 reliable, safe and effective prescription pain medications.

9 (b) That the guidelines appropriately limit both duration
10 and dosage of all prescribed pain medication drugs and include
11 consideration of treatment options beyond the prescription of
12 pain medication drugs.

13 (c) That the guidelines include measures to aid in the
14 management of all pain medications, specifically including
15 opioid medications.

16 (IV) The department shall receive, and shall at least
17 annually solicit through notice in the Pennsylvania Bulletin,
18 recommendations to revise its evidence-based prescription
19 guidelines for pain medications in furtherance of the factors in
20 subprovision (III). The department may revise these guidelines,
21 after consultation with the Department of Health and the
22 Department of Drug and Alcohol Programs and the appropriate
23 licensure boards under the Department of State. The department
24 shall publish notice of revisions to the guidelines in the
25 Pennsylvania Bulletin and the revisions shall take effect thirty
26 (30) days after the date of publication.

27 (V) The department shall ensure that these prescription
28 guidelines are available through the department's publicly
29 accessible Internet website for reference by physicians and the
30 general public.

1 (VI) The prescription of drugs consistent with or
2 recommended by the department's evidence-based prescription
3 guidelines shall be considered reasonable and necessary for the
4 purposes of this subclause. The prescription of drugs not
5 consistent with or recommended by the department's evidence-
6 based prescription guidelines shall nonetheless be considered
7 reasonable and necessary for the purposes of this paragraph if
8 the prescribing physician has submitted documentation explaining
9 the medical necessity of the prescription to the insurer or
10 self-insured employer at the time of the initial prescription on
11 a form prescribed by the department. An insurer or self-insured
12 employer shall accept such prescription unless the prescription
13 is determined not to be reasonable and necessary by a
14 utilization review organization in accordance with paragraph
15 (6).

16 (K) The department, in consultation with the Department of
17 Health and the Department of Drug and Alcohol Programs and the
18 appropriate licensing boards under the Department of State,
19 shall develop a program for educating and explaining its
20 evidence-based prescription guidelines on or before the
21 effective date of the prescription guidelines. All workers
22 compensation judges and all utilization review organizations
23 shall participate in this program within sixty (60) days of the
24 effective date of these prescription guidelines. The department
25 shall offer the program to all providers at locations and times
26 it deems appropriate. To the extent the program is presented in
27 written or video form, the department shall make it available
28 through the department's publicly accessible Internet website.

29 * * *

30 (6) Except in those cases in which a workers' compensation

1 judge asks for an opinion from peer review under section 420,
2 disputes as to reasonableness or necessity of treatment by a
3 health care provider shall be resolved in accordance with the
4 following provisions:

5 * * *

6 (ii) The department shall assign a request for utilization
7 review to a utilization review organization at random. The
8 utilization review organization shall issue a written report of
9 its findings and conclusions within the time frame required by
10 the nationally recognized accreditation standards adopted by the
11 department under subparagraph (v). In no case shall the report
12 of findings and conclusions be issued more than thirty (30) days
13 after the receipt of a request.

14 * * *

15 (v) The department shall approve only those utilization
16 review organizations that it determines have obtained
17 certification or accreditation by a nationally recognized
18 organization with certification or accreditation standards
19 appropriate for resolving utilization issues for workers'
20 compensation programs. The following shall apply:

21 (A) Within thirty (30) days of the effective date of this
22 subparagraph, the department shall publish notice in the
23 Pennsylvania Bulletin of the specific nationally recognized
24 certification or accreditation that will be required in order to
25 be approved as a utilization review organization.

26 (B) Upon publication under subclause (A), an entity without
27 the appropriate certification or accreditation may not engage in
28 utilization review under this act, except that an entity
29 approved as a utilization review organization by the department
30 prior to the effective date of this clause may continue to

1 engage in utilization review for up to eighteen (18) calendar
2 months after the publication of notice under subclause (A). If
3 the department determines that an entity approved as a
4 utilization review organization by the department prior to the
5 effective date of this subparagraph is actively attempting to
6 achieve the selected certification or accreditation, the entity
7 shall not be required to apply for reauthorization during the
8 eighteen-month period. A utilization review organization shall
9 adhere to the review standards of the selected nationally
10 recognized certification or accreditation organization for all
11 utilization review where the date of the injury is at least
12 eighteen (18) calendar months after the publication of notice
13 under subclause (A).

14 (C) The department shall enter an agreement with the
15 selected nationally recognized certification or accreditation
16 organization to provide for the certification or accreditation
17 process for utilization review organizations and employes of
18 utilization review organizations, including the costs of any
19 audits required for the certification or accreditation process.
20 The department shall make reasonable attempts to negotiate a
21 reduction of the cost of the certification or accreditation
22 process. An entity approved as a utilization review organization
23 by the department prior to the effective date of this
24 subparagraph, including a surviving association that results
25 from the merger of two or more utilization review organizations
26 under 15 Pa.C.S. Ch. 3 Subch. C (relating to merger), shall be
27 eligible to participate in the initial certification or
28 accreditation process at no cost to the entity. After January 1,
29 2020, an entity approved as a utilization review organization
30 shall be eligible to participate in the process to renew its

1 certification or accreditation at no cost to the entity. An
2 entity for which the department has incurred costs under this
3 subclause shall reimburse the department for its costs related
4 to the most recent certification or accreditation for the
5 entity, if the entity does not successfully obtain the initial
6 or renewal certification or accreditation. The actual amount of
7 the cost to the department for the certification or
8 accreditation process under this subclause, not to exceed one
9 million five hundred thousand dollars (\$1,500,000) annually,
10 shall be transferred to the department from the Workers'
11 Compensation Administration Fund.

12 (D) The department shall conduct outreach to all entities
13 approved as utilization review organizations by the department
14 prior to the effective date of this subparagraph. The outreach
15 shall include providing each entity with notice of the
16 requirements of this subparagraph, guidance on how this clause
17 will be enforced by the department and information on how the
18 entity may participate in the required certification or
19 accreditation process at no cost under subclause (C).

20 * * *

21 Section 2. Section 420 of the act is amended to read:

22 Section 420. (a) The board, the department or a workers'
23 compensation judge, if it or he deem it necessary, may, of its
24 or his own motion, either before, during, or after any hearing,
25 make or cause to be made an investigation of the facts set forth
26 in the petition or answer or facts pertinent in any injury under
27 this act. The board, department or workers' compensation judge
28 may appoint one or more impartial physicians or surgeons to
29 examine the injuries of the plaintiff and report thereon, or may
30 employ the services of such other experts as shall appear

1 necessary to ascertain the facts. The workers' compensation
2 judge when necessary or appropriate or upon request of a party
3 in order to rule on requests for review filed under section
4 306(f.1), or under other provisions of this act, may ask for an
5 opinion from peer review about the reasonableness or necessity
6 [or frequency] of treatment under section 306(f.1). The peer
7 review report or the peer report of any physician, surgeon, or
8 expert appointed by the department or by a workers' compensation
9 judge, including the report of a peer review organization, shall
10 be filed with the board or workers' compensation judge, as the
11 case may be, and shall be a part of the record and open to
12 inspection as such. The workers' compensation judge shall
13 consider the report as evidence but shall not be bound by such
14 report.

15 (b) The board or workers' compensation judge, as the case
16 may be, shall fix the compensation of such physicians, surgeons,
17 and experts, and other peer review organizations which, when so
18 fixed, shall be paid out of the Workmen's Compensation
19 Administration Fund.

20 (c) Peer review performed under this section and peer review
21 organizations used under this section shall comply with the
22 requirements established under section 306(f.1)(6).

23 Section 3. The Pennsylvania Compensation Rating Bureau shall
24 include in its annual loss cost filing to the Insurance
25 Department a calculation of any savings achieved through the
26 amendment of section 306(f.1) of the act and shall reduce the
27 filing to reflect those savings, subject to the Insurance
28 Department's review and approval.

29 Section 4. The Department of Labor and Industry shall submit
30 on or before January 1 of each year a report to the General

1 Assembly on the impact of the amendment of section 306(f.1) of
2 the act on the prescription of pain medications to injured
3 workers, including any recommendations for changes in the
4 regulation of those prescriptions.

5 Section 5. This act shall take effect in 60 days.