

**SENATE, No. 1102**

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**STATE OF NEW JERSEY**

**220th LEGISLATURE**

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INTRODUCED JANUARY 31, 2022

**Sponsored by:**  
**Senator JOSEPH F. VITALE**  
**District 19 (Middlesex)**

**SYNOPSIS**

Establishes limitations on and conditions associated with prescribers' acceptance of compensation from pharmaceutical manufacturers.

**CURRENT VERSION OF TEXT**

As introduced.



1    **AN ACT** concerning prescriber compensation by pharmaceutical  
2       manufacturers and supplementing Title 45 of the Revised  
3       Statutes.

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5       **BE IT ENACTED** *by the Senate and General Assembly of the State*  
6       *of New Jersey:*

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8       1. As used in this act:

9       “Bona fide services” means those services provided by a  
10      prescriber pursuant to an arrangement formalized in a written  
11      agreement including, but not limited to, presentations as speakers at  
12      promotional activities and continuing educational events,  
13      participation on advisory boards, and consulting arrangements. The  
14      written agreement shall specify the services to be provided, specify  
15      the dollar value of the consideration to be received by the prescriber  
16      based on the fair market value of the services, and identify the  
17      following: the legitimate need for services in advance; the  
18      connection between the competence, knowledge, and expertise of  
19      the prescriber and the purpose of the arrangement; how  
20      participation of the prescriber is reasonably related to achieving the  
21      identified purpose; the manner by which the prescriber will  
22      maintain records concerning the arrangement and the services  
23      provided by the prescriber; the venue and circumstances of any  
24      meeting in which the prescriber participates and how the venue and  
25      circumstances are conducive to the services provided and advance  
26      the primary focus of the meeting; and an attestation that the  
27      prescriber’s decision to render the services is not unduly influenced  
28      by a pharmaceutical manufacturer’s agent.

29      “Continuing education event” means a continuing education  
30      event, third-party scientific or educational conference, professional  
31      meeting, U.S. Food and Drug Administration required education  
32      and training, or any other gathering where responsibility for and  
33      control over the selection of content, faculty, educational methods,  
34      materials, and venue belongs to the event’s organizers in  
35      accordance with the standards of a nationally recognized  
36      accrediting entity, held in a venue that is appropriate and conducive  
37      to informational communication and training about healthcare  
38      information, where: the gathering is primarily dedicated, in both  
39      time and effort, to promoting objective scientific and educational  
40      activities and discourse, in which one or more educational  
41      presentations is the highlight of the gathering; and the main purpose  
42      for bringing attendees together is to further their knowledge on the  
43      topics being presented.

44      “Modest meals” means food or refreshment, the fair market  
45      value of which does not exceed \$15 for each prescriber, or such  
46      other amount as may be established by the Director of the Division  
47      of Consumer Affairs in the Department of Law and Public Safety by  
48      regulation.

1 “Non-faculty” means a prescriber who does not serve as a  
2 speaker or provide actual and substantive services as a faculty  
3 organizer or academic program consultant for a continuing  
4 education event or for a promotional activity.

5 “Pharmaceutical manufacturer” or “manufacturer” means any  
6 entity that: is engaged in the production, preparation, propagation,  
7 compounding, conversion, or processing of prescription drugs or  
8 biologics, by extraction from substances of natural origin or  
9 independently by means of chemical synthesis; or is directly  
10 engaged in the packaging, repackaging, labeling, relabeling, or  
11 distribution of prescription drugs or biologics. “Pharmaceutical  
12 manufacturer” or “manufacturer” does not include a health care  
13 facility licensed by the Department of Health or a pharmacy holding  
14 a permit issued by the New Jersey State Board of Pharmacy.

15 “Pharmaceutical manufacturer’s agent” or “manufacturer’s  
16 agent” means a person who, while employed by or under contract  
17 with a pharmaceutical manufacturer, engages in detailing,  
18 promotional activities, or other marketing of prescription drugs or  
19 biologics to: a prescriber authorized to prescribe, dispense, or  
20 purchase prescription drugs or biologics; a health care facility; or a  
21 pharmacist. “Pharmaceutical manufacturer’s agent” or  
22 “manufacturer’s agent” does not include a prescriber or pharmacist  
23 when acting within the ordinary scope of the practice for which the  
24 prescriber or pharmacist is licensed.

25 “Prescriber” means a physician, podiatrist, physician assistant,  
26 advanced practice nurse, dentist, or optometrist licensed pursuant to  
27 Title 45 of the Revised Statutes. “Prescriber” does not include a  
28 licensee who is an employee of a pharmaceutical manufacturer who  
29 does not provide patient care.

30 “Promotional activity” means any unaccredited activity, meeting,  
31 or program organized or sponsored by a pharmaceutical  
32 manufacturer or the manufacturer’s agent that is directed at  
33 prescribers to promote the prescription, recommendation, supply,  
34 administration, use, or consumption of the manufacturer’s products  
35 through any medium.

36  
37 2. a. A prescriber shall not accept, directly or indirectly, any  
38 financial benefit or benefit-in-kind, including, but not limited to,  
39 gifts, payments, stock, stock options, grants, scholarships,  
40 subsidies, or charitable contributions, except as permitted under  
41 section 3 of this act, from any manufacturer or manufacturer’s  
42 agent.

43 b. A prescriber shall not accept, directly or indirectly, any  
44 entertainment or recreational items, such as tickets to theater or  
45 sporting events, or leisure or vacation trips, from any manufacturer  
46 or manufacturer’s agent.

47 c. Except as permitted under section 3 of this act, a prescriber  
48 shall not accept from any manufacturer or manufacturer’s agent any

1 item of value that does not advance disease or treatment education,  
2 including:

3 (1) pens, note pads, clipboards, mugs, or other items with a  
4 company or product logo;

5 (2) items intended for the personal benefit of the prescriber or  
6 staff, such as floral arrangements, sporting equipment, artwork, or  
7 items that may have utility in both the professional and non-  
8 professional setting, such as electronic devices;

9 (3) any payment in cash or cash equivalent, such as a gift card  
10 or gift certificate; or

11 (4) any payment or direct subsidy to a non-faculty prescriber to  
12 support attendance at, or as remuneration for time spent attending,  
13 or for the costs of travel, lodging, or other personal expenses  
14 associated with attending, any continuing education event or  
15 promotional activity.

16 d. A prescriber shall not accept meals from any manufacturer  
17 or manufacturer's agent, except as provided in section 3 of this act.

18 e. Unless an immediate family member is employed by a  
19 manufacturer and receives, as part of the usual and customary  
20 employment relationship, compensation, financial benefit, or other  
21 item of value, the prohibitions listed in this rule shall also apply to  
22 the prescriber's immediate family. For purposes of this subsection,  
23 "immediate family" means an individual's spouse, civil union  
24 partner, or domestic partner, or the individual's or spouse's, civil  
25 union partner's, or domestic partner's parent, child, brother, sister,  
26 aunt, uncle, niece, nephew, grandparent, grandchild, son-in-law,  
27 daughter-in-law, stepparent, stepchild, stepbrother, stepsister, half-  
28 brother, or half-sister, whether the relative is related to the  
29 individual or the individual's spouse, civil union partner, or  
30 domestic partner by blood, marriage, or adoption.

31

32 3. Consistent with the requirements of this section, a prescriber  
33 may accept the following from a manufacturer or manufacturer's  
34 agent:

35 a. Items designed primarily for educational purposes for the  
36 prescriber or patients that have minimal or no value to the prescriber  
37 outside of the prescriber's professional responsibilities. Examples of  
38 educational items include anatomical models for use in an examination  
39 room or other information and materials in any form directly related to  
40 patient care or prescriber education. Items that may have an  
41 independent value to the prescriber outside of the prescriber's  
42 professional responsibilities, such as electronic devices, may only be  
43 accepted if they are used by patients and remain in a common area of  
44 the prescriber's office.

45 b. A manufacturer-subsidized registration fee at a continuing  
46 education event if that fee is available to all event participants.

47 c. Modest meals provided through the event organizer at a  
48 continuing education event, provided the meals facilitate the  
49 educational program to maximize prescriber learning.

1 d. Modest meals provided to non-faculty prescribers through  
2 promotional activities no more than four times in a calendar year from  
3 the same manufacturer.

4 e. Compensation, based on fair market value, for providing bona  
5 fide services as a speaker or faculty organizer or academic program  
6 consultant for a continuing education event. A prescriber serving in  
7 this capacity also may accept reasonable payment and remuneration  
8 for travel, lodging, and other personal expenses associated with such  
9 services. A prescriber may be granted continuing education credit for  
10 participation in such activities if the continuing education  
11 requirements of the prescriber's professional licensing board are  
12 satisfied.

13 f. Compensation, based on fair market value, for providing bona  
14 fide services as a speaker or faculty organizer or academic program  
15 consultant for a promotional activity, consistent with the limits set  
16 forth in section 5 of this act. A prescriber serving in this capacity may  
17 also accept reasonable payment or remuneration for travel, lodging,  
18 and other personal expenses associated with such services. A  
19 prescriber may not claim continuing education credit for participation  
20 in such activities.

21 g. Compensation, based on fair market value, for participation on  
22 advisory bodies or under consulting arrangements, consistent with the  
23 limits set forth in section 5 of this act.

24  
25 4. A prescriber may accept sample medications or devices that  
26 are intended to be used exclusively for the benefit of the  
27 prescriber's patients, provided the prescriber does not charge  
28 patients for such samples, and all applicable dispensing standards  
29 set forth in the prescriber's licensing board rules are satisfied.

30  
31 5. A prescriber shall not accept more than \$10,000, or such  
32 other amount as may be established by the Director of the Division  
33 of Consumer Affairs in the Department of Law and Public Safety by  
34 regulation, in the aggregate from all manufacturers in any calendar  
35 year for the bona fide services of presentations as a speaker or  
36 faculty organizer or academic program consultant at promotional  
37 activities, participation on advisory boards, and consulting  
38 arrangements. Payments for speaking at continuing education  
39 events are not subject to this limit, but shall be in an amount that  
40 constitutes fair market value and is set forth in a written agreement.

41  
42 6. A prescriber serving as a speaker at a continuing education  
43 event or for a promotional activity shall directly disclose to  
44 attendees either orally or in writing at the beginning of the  
45 presentation whether the prescriber has accepted payment for bona  
46 fide services from the sponsoring manufacturer within the preceding  
47 five years.

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1        7. A prescriber who is employed by a manufacturer and who  
2        also provides patient care shall comply with the disclosure  
3        requirements of section 6 of this act, but is exempt from the  
4        compensation prohibitions of this act.

6        8. The Director of the Division of Consumer Affairs in the  
7 Department of Law and Public Safety, in accordance with the  
8 “Administrative Procedure Act,” P.L.1968, c.410 (C.52:14B-  
9 1 et seq.), shall adopt any rules and regulations as the director  
10 deems necessary to carry out the provisions of this act.

9. This act shall take effect on the first day of the seventh month next following the date of enactment, except that the Director of the Division of Consumer Affairs in the Department of Law and Public Safety may take any anticipatory administrative action in advance as shall be necessary for the implementation of this act.

## STATEMENT

22 This bill establishes limitations on, and conditions associated  
23 with, prescribers' acceptance of compensation from pharmaceutical  
24 manufacturers.

Licensed physicians, podiatrists, physician assistants, advanced practice nurses, dentists, and optometrists are all authorized to prescribe pharmaceutical products within the scope of their professional practice. These professionals are also permitted to enter into financial relationships with drug manufacturers, which are limited in varying ways by the different licensing boards, but which may raise concerns that these relationships might influence prescriber's treatment decisions to the detriment of their patients.

The bill generally prohibits prescribers and members of their immediate families from accepting, from a pharmaceutical manufacturer or a manufacturer's agent, any financial benefits or benefits-in-kind, entertainment or recreational items, things of value that do not advance disease or treatment education, or meals. A prescriber would be permitted to accept gifts or payments that meet certain conditions, including: items that are used primarily for educational purposes for the prescriber or patients that have minimal value outside of the prescriber's professional responsibilities; subsidized registration fees at continuing education events if that subsidized fee is available to all participants; modest meals provided at a continuing education event; up to four modest meals at promotional activities per year; and compensation based on fair market value for bona fide services as a speaker or organizer or consultant at a continuing education event.

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1       The bill permits prescribers to accept sample medications and  
2 devices exclusively for the benefit of the prescriber's patients,  
3 provided that the prescriber does not charge patients for such  
4 samples and all requirements of the prescriber's licensing board are  
5 met.

6       The bill also permits a prescriber to receive compensation for  
7 bona fide services as a speaker or faculty organizer or academic  
8 program consultant at promotional events, participation on advisory  
9 boards, and other consulting arrangements, subject to a cap of \$10,000  
10 per year in aggregate. The \$10,000 cap may be altered by the  
11 Director of the Division of Consumer Affairs in the Department of  
12 Law and Public Safety by regulation.

13       Under the bill, a prescriber who serves as a speaker at a continuing  
14 education or promotional event would be required to disclose whether  
15 the prescriber has accepted payment for bona fide services from the  
16 sponsoring manufacturer in the preceding five years.

17       A prescriber employed by a manufacturer who also provides patient  
18 care would be subject to the disclosure requirements of the bill, but  
19 exempt from the compensation provisions.