

# Senate Bill 543

Sponsored by Senator MONNES ANDERSON

## SUMMARY

The following summary is not prepared by the sponsors of the measure and is not a part of the body thereof subject to consideration by the Legislative Assembly. It is an editor's brief statement of the essential features of the measure **as introduced**.

Creates method by which health care practitioner may offer to treat patient who has terminal disease with drug, biological product or device not approved by United States Food and Drug Administration. Provides protections, including waiver of liability, for health care practitioners, health care facilities and professional organizations or associations that comply with Act.

## A BILL FOR AN ACT

1  
2 Relating to treatments for patients with terminal diseases.

3 **Be It Enacted by the People of the State of Oregon:**

4 **SECTION 1. As used in sections 1 to 12 of this 2015 Act:**

5 (1) "Attending physician" means the physician who has primary responsibility for the  
6 care of a patient.

7 (2) "Capable" means that, in the opinion of an attending physician, consulting physician  
8 or other health care practitioner, a patient has the ability to make and communicate health  
9 care decisions to health care practitioners, including the ability to communicate through  
10 individuals familiar with the patient's manner of communicating.

11 (3) "Consulting physician" means a physician who is qualified by specialty or experience  
12 to diagnose a patient who has a terminal disease and to make a prognosis for that patient.

13 (4) "Health care facility" has the meaning given that term in ORS 442.015.

14 (5) "Health care practitioner" means an individual who is licensed, certified or otherwise  
15 authorized by the laws of this state to provide health care services or to dispense drugs or  
16 biological products.

17 (6) "Physician" means a doctor of medicine or osteopathy licensed to practice medicine  
18 under ORS chapter 677.

19 (7) "Terminal disease" means an irreversible disease that:

20 (a) Is not treatable by drugs, biological products or devices approved by the United States  
21 Food and Drug Administration; and

22 (b) In a physician's reasonable medical judgment, will result in the patient's death within  
23 one year.

24 **SECTION 2. (1) The attending physician of a patient who has a terminal disease may re-**  
25 **fer the patient to a health care practitioner who offers treatment described in section 3 of**  
26 **this 2015 Act if:**

27 (a) The treatment is being offered only for purposes related to the terminal disease;

28 (b) The patient is capable, 18 years of age or older and a resident of this state; and

29 (c) The attending physician informs the patient:

30 (A) That the patient has a terminal disease;

**NOTE:** Matter in **boldfaced** type in an amended section is new; matter [*italic and bracketed*] is existing law to be omitted. New sections are in **boldfaced** type.

1 (B) Of the attending physician's prognosis for the patient;

2 (C) That the drug, biological product or device to be used in treating the patient is not  
3 approved by the United States Food and Drug Administration;

4 (D) Of each potential risk associated with receiving the treatment that is known to the  
5 attending physician; and

6 (E) Of feasible alternatives to receiving the treatment, including palliative care, hospice  
7 care and pain control.

8 (2) A patient who has a terminal disease may demonstrate the patient's Oregon residency  
9 to the patient's attending physician by presenting:

10 (a) A driver license, driver permit or identification card issued to the patient by the De-  
11 partment of Transportation;

12 (b) Evidence that the patient is registered to vote in this state;

13 (c) Evidence that the patient owns or leases property in this state; or

14 (d) A copy of the patient's Oregon individual tax return for the immediately preceding  
15 tax year.

16 **SECTION 3.** (1) A health care practitioner may offer to treat a patient who has a ter-  
17 minal disease with a drug, biological product or device not approved by the United States  
18 Food and Drug Administration only if:

19 (a) The health care practitioner is authorized by the laws of this state to provide health  
20 care services or to dispense drugs or biological products, and the health care practitioner is  
21 acting within the scope of that authority;

22 (b) The drug, biological product or device has undergone Phase I of a clinical trial, as-  
23 sessing the safety of the drug, biological product or device;

24 (c) The treatment is provided free of charge to the patient;

25 (d) The treatment is being offered only for purposes related to the terminal disease;

26 (e) The patient is capable, 18 years of age or older and a resident of this state;

27 (f) The patient was referred to the health care practitioner by the patient's attending  
28 physician under section 2 of this 2015 Act; and

29 (g) The health care practitioner refers the patient to a consulting physician to affirm the  
30 attending physician's diagnosis and prognosis.

31 (2) A patient who has a terminal disease may demonstrate the patient's Oregon residency  
32 to the health care practitioner by presenting:

33 (a) A driver license, driver permit or identification card issued to the patient by the De-  
34 partment of Transportation;

35 (b) Evidence that the patient is registered to vote in this state;

36 (c) Evidence that the patient owns or leases property in this state; or

37 (d) A copy of the patient's Oregon individual tax return for the immediately preceding  
38 tax year.

39 **SECTION 4.** Before a patient may receive treatment described in section 3 of this 2015  
40 Act, a consulting physician must examine the patient and confirm, in writing:

41 (1) The attending physician's diagnosis that the patient has a terminal disease;

42 (2) The attending physician's prognosis for the patient;

43 (3) That the patient is capable; and

44 (4) That the patient knows:

45 (a) That the drug, biological product or device to be used in treating the patient is not

1 approved by the United States Food and Drug Administration;

2 (b) Of each potential risk associated with receiving the treatment known to the consult-  
3 ing physician; and

4 (c) Of feasible alternatives to receiving the treatment, including palliative care, hospice  
5 care and pain control.

6 **SECTION 5.** Upon receiving an offer for treatment described in section 3 of this 2015 Act,  
7 a patient who has a terminal disease who is capable, 18 years of age or older and a resident  
8 of this state may elect to receive that treatment by signing and dating a form attesting to  
9 the election in the presence of two witnesses. A form attesting to an election must include:

10 (1) The attending physician's diagnosis for the patient;

11 (2) The attending physician's prognosis for the patient;

12 (3) A statement that the drug, biological product or device to be used in treating the  
13 patient is not approved by the United States Food and Drug Administration;

14 (4) A description of each potential risk that is associated with receiving the treatment;

15 (5) A waiver of liability for any act or omission of an act related to administering the  
16 treatment that does not constitute gross negligence for:

17 (a) Any health care practitioner who participates in administering the treatment; or

18 (b) Any health care facility or professional organization or association involved in the  
19 administration of the treatment;

20 (6) A provision authorizing any information obtained during the treatment to be used:

21 (a) By the inventor, manufacturer or supplier of any drug, biological product or device  
22 used in treating the patient for research, analytical or marketing purposes; and

23 (b) By any health care practitioner who participates in administering the treatment for  
24 research or analytical purposes; and

25 (7) A statement signed and dated by both witnesses attesting that the patient, to the best  
26 of the witnesses' knowledge, is capable and acting voluntarily.

27 **SECTION 6.** (1) Of the witnesses described in section 5 of this 2015 Act, one must be an  
28 individual who is not:

29 (a) A relative of the patient by blood, marriage or adoption;

30 (b) A person who, at the time the form is signed, would be entitled to any portion of the  
31 estate of the patient upon the patient's death under any will or by operation of law; or

32 (c) An owner, operator or employee of a health care facility where the patient resides  
33 or receives health care services.

34 (2) Neither witness described in section 5 of this 2015 Act may be the attending physician  
35 of the patient.

36 **SECTION 7.** A waiver of liability required by section 5 (5) of this 2015 Act must be written  
37 in plain and simple language.

38 **SECTION 8.** (1) Except as provided in subsection (2) of this section, a health care prac-  
39 titioner who participates in administering a treatment described in section 3 of this 2015 Act,  
40 or a health care facility or professional organization or association involved in the adminis-  
41 tration of the treatment, is not subject to civil or criminal liability for acts or omissions of  
42 acts related to administering the treatment if the administration of the treatment complies  
43 with sections 1 to 12 of this 2015 Act.

44 (2) This section does not apply to acts or omissions of acts that constitute gross  
45 negligence.

1       **SECTION 9.** (1) Except as provided in subsection (2) of this section and sections 10 and  
2 11 of this 2015 Act, a licensing board, health care facility, health care practitioner or pro-  
3 fessional organization or association may not subject a health care practitioner to discipline,  
4 including suspension, loss of license, loss of privileges, loss of membership or any other  
5 penalty, for participating in administering a treatment described in section 3 of this 2015 Act  
6 if the administration of the treatment complies with sections 1 to 12 of this 2015 Act.

7       (2) This section does not apply to acts or omissions of acts that constitute gross  
8 negligence.

9       **SECTION 10.** A health care facility or health care practitioner may prohibit another  
10 health care practitioner from participating in administering a treatment described in section  
11 3 of this 2015 Act at the health care facility, or on premises owned or controlled by the  
12 prohibiting health care practitioner, if the health care facility or prohibiting health care  
13 practitioner:

14       (1) Adopts a written policy that clearly prohibits the administration of such treatment;  
15 and

16       (2) Provides a printed or electronic copy of that policy to each health care practitioner  
17 who provides health care services at the health care facility or on premises owned or con-  
18 trolled by the prohibiting health care practitioner.

19       **SECTION 11.** If a health care practitioner violates a prohibition authorized by section 10  
20 of this 2015 Act:

21       (1) A licensing board, health care facility, health care practitioner or professional or-  
22 ganization or association may impose upon the violating health care practitioner any form  
23 of discipline described in section 9 of this 2015 Act that the licensing board, health care fa-  
24 cility, health care practitioner or professional organization or association otherwise may le-  
25 gally impose; and

26       (2) The health care facility or prohibiting health care practitioner may:

27       (a) Terminate any lease or other property contract entered into with the violating health  
28 care practitioner and subject the violating health care practitioner to any other nonmonetary  
29 remedies provided by such a contract; or

30       (b) Terminate any contract for the provision of services entered into with the violating  
31 health care practitioner and subject the violating health care practitioner to any other non-  
32 monetary remedies provided by such a contract.

33       **SECTION 12.** Sections 1 to 12 of this 2015 Act do not require an insurer to reimburse any  
34 cost associated with treatment described in section 3 of this 2015 Act.

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