

**PROPOSED AMENDMENTS TO  
A-ENGROSSED HOUSE BILL 2300**

1 On page 1 of the printed A-engrossed bill, delete lines 4 through 26 and  
2 delete pages 2 through 5 and insert:

3 **“SECTION 1. As used in sections 1 to 14 of this 2015 Act:**

4 **“(1) ‘Attending physician’ means the physician who has primary**  
5 **responsibility for the care of a patient.**

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

11 **“(3) ‘Consulting physician’ means a physician who is qualified by**  
12 **specialty or experience to diagnose a patient who has a terminal dis-**  
13 **ease and to make a prognosis for that patient.**

14 **“(4) ‘Health care facility’ has the meaning given that term in ORS**  
15 **442.015.**

16 **“(5) ‘Health care practitioner’ means an individual who is licensed,**  
17 **certified or otherwise authorized by the laws of this state to provide**  
18 **health care services or to dispense drugs.**

19 **“(6) ‘Investigational product’ means a drug, biological product or**  
20 **device that has successfully completed Phase I and is currently in**  
21 **Phase II or a subsequent phase of an approved clinical trial, as defined**  
22 **in ORS 743A.192, assessing the safety of the drug, biological product**

1 or device.

2 “(7) ‘Physician’ means a doctor of medicine or osteopathy licensed  
3 to practice medicine under ORS chapter 677.

4 “(8) ‘Qualified’ means, with respect to a patient, that the patient  
5 is:

6 “(a) Capable;

7 “(b) A resident of this state; and

8 “(c) 18 years of age or older.

9 “(9) ‘Terminal disease’ means an illness or a medical or surgical  
10 condition that in a physician’s reasonable medical judgment will result  
11 in the patient’s death within six months.

12 “SECTION 2. (1) The attending physician of a patient who has a  
13 terminal disease may refer the patient to a health care practitioner  
14 who offers treatment as described in section 3 of this 2015 Act if:

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

17 “(b) The patient is qualified;

18 “(c) In the attending physician’s judgment, the patient is acting  
19 voluntarily and is not being coerced; and

20 “(d) The attending physician informs the patient:

21 “(A) That the patient has a terminal disease;

22 “(B) Of the attending physician’s prognosis for the patient;

23 “(C) That the investigational product to be used in treating the  
24 patient is not approved by the United States Food and Drug Adminis-  
25 tration and that the investigational product may not be effective in  
26 treating the patient;

27 “(D) Of each potential risk associated with receiving the treatment  
28 that is known to the attending physician;

29 “(E) That to receive the treatment, the patient may be required to  
30 pay the costs of administering the treatment and the costs of, or the

1 costs associated with, manufacturing the investigational product as  
2 described in section 3 (1)(b) of this 2015 Act;

3 “(F) That to receive the treatment, the patient must waive liability  
4 as described in section 5 (5) of this 2015 Act;

5 “(G) That receiving the treatment relieves an insurer of reimburs-  
6 ing costs as described in section 12 of this 2015 Act;

7 “(H) Of feasible alternatives to receiving the treatment, including  
8 palliative care, hospice care and pain control; and

9 “(I) That expanded access to treating the patient’s terminal disease  
10 may be provided pursuant to 21 C.F.R. 312.300 to 312.320 and may be  
11 an option for the patient, and, depending on the type of coverage the  
12 patient’s insurer provides, that a patient might not be required to pay  
13 the costs of administering a treatment provided pursuant to 21 C.F.R.  
14 312.300 to 312.320, or the costs of, or the costs associated with, manu-  
15 facturing an investigational product used to treat a patient pursuant  
16 to 21 C.F.R. 312.300 to 312.320.

17 “(2) A patient who has a terminal disease may demonstrate the  
18 patient’s Oregon residency to the patient’s attending physician by  
19 presenting:

20 “(a) A driver license, driver permit or identification card issued to  
21 the patient by the Department of Transportation;

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

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

24 or

25 “(d) A copy of the patient’s Oregon individual tax return for the  
26 immediately preceding tax year.

27 “(3) If in the opinion of an attending physician a patient is suffering  
28 from a psychiatric or psychological disorder or depression causing  
29 impaired judgment, the attending physician shall refer the patient for  
30 counseling. Treatment may not be provided as described in section 3

1 of this 2015 Act until the person performing the counseling determines  
2 that the patient is not suffering from a psychiatric or psychological  
3 disorder or depression causing impaired judgment.

4 **“SECTION 3. (1) A health care practitioner may offer to treat a**  
5 **patient who has a terminal disease with an investigational product not**  
6 **approved by the United States Food and Drug Administration only if:**

7 **“(a) The health care practitioner is authorized by the laws of this**  
8 **state to provide health care services or to dispense drugs, and the**  
9 **health care practitioner is acting within the scope of that authority;**

10 **“(b) The treatment is provided to the patient for no more than the**  
11 **costs of administering the treatment and the costs of, or the costs**  
12 **associated with, manufacturing the investigational product;**

13 **“(c) The patient is not compensated for receiving the treatment;**

14 **“(d) The treatment is being offered only for purposes related to the**  
15 **terminal disease;**

16 **“(e) The patient is qualified;**

17 **“(f) The patient was referred to the health care practitioner by the**  
18 **patient’s attending physician under section 2 of this 2015 Act;**

19 **“(g) The health care practitioner refers the patient to a consulting**  
20 **physician to confirm the attending physician’s diagnosis and**  
21 **prognosis; and**

22 **“(h) In the health care practitioner’s judgment, the patient is act-**  
23 **ing voluntarily and is not being coerced.**

24 **“(2) A patient who has a terminal disease may demonstrate the**  
25 **patient’s Oregon residency to the health care practitioner by present-**  
26 **ing:**

27 **“(a) A driver license, driver permit or identification card issued to**  
28 **the patient by the Department of Transportation;**

29 **“(b) Evidence that the patient is registered to vote in this state;**

30 **“(c) Evidence that the patient owns or leases property in this state;**

1 or

2 “(d) A copy of the patient’s Oregon individual tax return for the  
3 immediately preceding tax year.

4 “(3) If in the opinion of the health care practitioner a patient is  
5 suffering from a psychiatric or psychological disorder or depression  
6 causing impaired judgment, the health care practitioner shall refer the  
7 patient for counseling. Treatment may not be provided as described in  
8 this section until the person performing the counseling determines  
9 that the patient is not suffering from a psychiatric or psychological  
10 disorder or depression causing impaired judgment.

11 “(4) If a patient accepts an offer for treatment under this section,  
12 and if the patient has health insurance, the health care practitioner  
13 offering to treat the patient must notify the insurer that the patient  
14 is receiving the treatment.

15 “SECTION 4. (1) Before a patient may receive treatment as de-  
16 scribed in section 3 of this 2015 Act, a consulting physician must ex-  
17 amine the patient and confirm, in writing:

18 “(a) The attending physician’s diagnosis that the patient has a ter-  
19 minal disease;

20 “(b) The attending physician’s prognosis for the patient;

21 “(c) That the patient is qualified;

22 “(d) That in the consulting physician’s judgment the patient is  
23 acting voluntarily and is not being coerced; and

24 “(e) That the patient is informed:

25 “(A) That the investigational product to be used in treating the  
26 patient is not approved by the United States Food and Drug Adminis-  
27 tration and that the investigational product may not be effective in  
28 treating the patient;

29 “(B) Of each potential risk associated with receiving the treatment  
30 known to the consulting physician;

1       **“(C) That to receive the treatment, the patient may be required to**  
2 **pay the costs of administering the treatment and the costs of, or the**  
3 **costs associated with, manufacturing the investigational product as**  
4 **described in section 3 (1)(b) of this 2015 Act;**

5       **“(D) That to receive the treatment, the patient must waive liability**  
6 **as described in section 5 (5) of this 2015 Act;**

7       **“(E) That receiving the treatment relieves an insurer of reimburs-**  
8 **ing costs as described in section 12 of this 2015 Act;**

9       **“(F) Of feasible alternatives to receiving the treatment, including**  
10 **palliative care, hospice care and pain control; and**

11       **“(G) That expanded access to treating the patient’s terminal disease**  
12 **may be provided pursuant to 21 C.F.R. 312.300 to 312.320 and may be**  
13 **an option for the patient, and, depending on the type of coverage the**  
14 **patient’s insurer provides, that a patient might not be required to pay**  
15 **the costs of administering a treatment provided pursuant to 21 C.F.R.**  
16 **312.300 to 312.320, or the costs of, or the costs associated with, manu-**  
17 **facturing an investigational product used to treat a patient pursuant**  
18 **to 21 C.F.R. 312.300 to 312.320.**

19       **“(2) A patient who has a terminal disease may demonstrate the**  
20 **patient’s Oregon residency to the consulting physician by presenting:**

21       **“(a) A driver license, driver permit or identification card issued to**  
22 **the patient by the Department of Transportation;**

23       **“(b) Evidence that the patient is registered to vote in this state;**

24       **“(c) Evidence that the patient owns or leases property in this state;**

25 **or**

26       **“(d) A copy of the patient’s Oregon individual tax return for the**  
27 **immediately preceding tax year.**

28       **“(3) If in the opinion of the consulting physician a patient is suf-**  
29 **fering from a psychiatric or psychological disorder or depression**  
30 **causing impaired judgment, the consulting physician shall refer the**

1 patient for counseling. Treatment may not be provided as described in  
2 section 3 of this 2015 Act until the person performing the counseling  
3 determines that the patient is not suffering from a psychiatric or  
4 psychological disorder or depression causing impaired judgment.

5 **“SECTION 5. Upon receiving an offer for treatment as described in**  
6 **section 3 of this 2015 Act, a patient who has a terminal disease and**  
7 **who is qualified may elect to receive that treatment by signing and**  
8 **dating a form attesting to the election in the presence of two wit-**  
9 **nesses. 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 investigational product to be used in**  
13 **treating the patient is not approved by the United States Food and**  
14 **Drug Administration;**

15 **“(4) A description of each potential risk that is associated with re-**  
16 **ceiving the treatment;**

17 **“(5) A waiver of liability for any act or omission of an act related**  
18 **to administering the treatment or manufacturing or distributing the**  
19 **investigational product that does not constitute gross negligence for:**

20 **“(a) Any health care practitioner who participates in administering**  
21 **the treatment, to whom a health care practitioner who participates in**  
22 **administering the treatment refers the patient or with whom a health**  
23 **care practitioner who participates in administering the treatment**  
24 **consults;**

25 **“(b) Any health care facility or professional organization or associ-**  
26 **ation involved in the administration of the treatment; or**

27 **“(c) Any person that participates in manufacturing or distributing**  
28 **the investigational product used to treat the patient;**

29 **“(6) A provision authorizing any information obtained during the**  
30 **treatment to be used:**

1       “(a) By the inventor, manufacturer or supplier of any  
2       investigational product used in treating the patient for research, ana-  
3       lytical or marketing purposes; and

4       “(b) By any health care practitioner who participates in adminis-  
5       tering the treatment for research or analytical purposes; and

6       “(7) A statement signed and dated by both witnesses attesting that  
7       the patient, to the best of the witnesses’ knowledge, is capable and  
8       acting voluntarily.

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

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

12      “(b) An individual who, at the time the form is signed, would be  
13      entitled to any portion of the estate of the patient upon the patient’s  
14      death under any will or by operation of law; or

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

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

19      “SECTION 7. A waiver of liability required by section 5 (5) of this  
20      2015 Act must be written in plain and simple language.

21      “SECTION 8. (1) Except as provided in subsection (3) of this section,  
22      a health care practitioner who participates in administering a treat-  
23      ment as described in section 3 of this 2015 Act, or a health care facility  
24      or professional organization or association involved in the adminis-  
25      tration of the treatment, is not subject to civil or criminal liability for  
26      acts or omissions of acts related to administering the treatment if the  
27      administration of the treatment complies with sections 1 to 14 of this  
28      2015 Act.

29      “(2) Except as provided in subsection (3) of this section, a man-  
30      ufacturer or distributor of an investigational product used to treat a

1 patient pursuant to section 3 of this 2015 Act is not subject to civil or  
2 criminal liability for acts or omissions of acts related to the adminis-  
3 tration of the investigational product.

4 “(3) This section does not apply to acts or omissions of acts that  
5 constitute gross negligence.

6 “SECTION 9. (1) Except as provided in subsection (2) of this section  
7 and sections 10 and 11 of this 2015 Act, a licensing board, health care  
8 facility, health care practitioner or professional organization or asso-  
9 ciation may not subject a health care practitioner to discipline, in-  
10 cluding suspension, loss of license, loss of privileges, loss of  
11 membership or any other penalty, for participating in administering  
12 a treatment as described in section 3 of this 2015 Act if the adminis-  
13 tration of the treatment complies with sections 1 to 14 of this 2015 Act.

14 “(2) This section does not apply to acts or omissions of acts that  
15 constitute gross negligence.

16 “SECTION 10. A health care facility or health care practitioner may  
17 prohibit another health care practitioner from participating in ad-  
18 ministering a treatment as described in section 3 of this 2015 Act at  
19 the health care facility or on premises owned or controlled by the  
20 prohibiting health care practitioner.

21 “SECTION 11. If a health care practitioner violates a prohibition  
22 authorized by section 10 of this 2015 Act:

23 “(1) A licensing board, health care facility, health care practitioner  
24 or professional organization or association may impose upon the vio-  
25 lating health care practitioner any form of discipline referred to in  
26 section 9 of this 2015 Act that the licensing board, health care facility,  
27 health care practitioner or professional organization or association  
28 otherwise may legally impose; and

29 “(2) The health care facility or prohibiting health care practitioner  
30 may:

1       “(a) Terminate any lease or other property contract entered into  
2 with the violating health care practitioner and subject the violating  
3 health care practitioner to any other nonmonetary remedies provided  
4 by such a contract; or

5       “(b) Terminate any contract for the provision of services entered  
6 into with the violating health care practitioner and subject the vio-  
7 lating health care practitioner to any other nonmonetary remedies  
8 provided by such a contract.

9       “SECTION 12. Sections 1 to 14 of this 2015 Act do not require an  
10 insurer to reimburse any cost:

11       “(1) Associated with undergoing a treatment as described in section  
12 3 of this 2015 Act; or

13       “(2) Demonstrated to be associated with an adverse effect that is a  
14 result of undergoing a treatment as described in section 3 of this 2015  
15 Act.

16       “SECTION 13. Eligibility for hospice care must be determined on  
17 the basis of a patient’s overall prognosis and care or treatment goals  
18 as determined by the patient’s attending physician and may not be  
19 determined on the basis of whether a patient is undergoing or has  
20 undergone a treatment as described in section 3 of this 2015 Act.

21       “SECTION 14. (1) The Oregon Health Authority shall annually re-  
22 view a sample of records maintained pursuant to sections 1 to 14 of  
23 this 2015 Act.

24       “(2) An attending physician who makes a referral under section 2  
25 of this 2015 Act, a health care practitioner who administers treatment  
26 as described in section 3 of this 2015 Act and a consulting physician  
27 who provides written confirmation as described in section 4 of this 2015  
28 Act must file with the authority a record, in a form and manner pre-  
29 scribed by the authority, of the findings of the attending physician,  
30 health care practitioner or consulting physician.

1       **“(3) At a minimum, the authority shall require that a record filed**  
2 **by a health care practitioner who administers treatment as described**  
3 **in section 3 of this 2015 Act must include:**

4       **“(a) The adverse effects of the treatment, if any;**

5       **“(b) The positive outcomes of the treatment, if any;**

6       **“(c) The cost of the treatment to the patient; and**

7       **“(d) The demographics of the patients to whom the treatment is**  
8 **administered.**

9       **“(4) The authority shall adopt rules to facilitate the collection of**  
10 **information required to comply with sections 1 to 14 of this 2015 Act,**  
11 **including rules related to the submission of information required by**  
12 **this section. Except as otherwise provided by law, information col-**  
13 **lected by the authority under this section is not a public record and**  
14 **is not available for inspection by the public.**

15       **“(5) The authority shall generate and make available to the public**  
16 **an annual statistical report of information collected by the authority**  
17 **pursuant to this section and of patients who receive treatment pro-**  
18 **vided pursuant to 21 C.F.R. 312.300 to 312.320.**

19       **“(6) The authority shall make the annual report generated under**  
20 **subsection (5) of this section available to the Legislative Assembly, in**  
21 **the manner required by ORS 192.245, on or before February 1 of each**  
22 **odd-numbered year.**

23       **“SECTION 15. This 2015 Act is repealed on January 2, 2022.”.**

24