

**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 that:

12 “(i) A patient who receives treatment pursuant to 21 C.F.R. 312.300
13 to 312.320 is not required to pay the costs of administering the treat-
14 ment or the costs of, or the costs associated with, manufacturing the
15 investigational product as described in section 3 (1)(b) of this 2015 Act;
16 and

17 “(ii) An insurer is not relieved of reimbursing costs as described in
18 section 12 of this 2015 Act for treatment provided pursuant to 21 C.F.R.
19 312.300 to 312.320.

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

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

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

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

27 or

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

30 “(3) If in the opinion of an attending physician a patient is suffering

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

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

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

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

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

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

19 **“(e) The patient is qualified;**

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

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

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

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

30 **“(a) A driver license, driver permit or identification card issued to**

1 the patient by the Department of Transportation;

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

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

4 or

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

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

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

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

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

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

24 “(c) That the patient is qualified;

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

27 “(e) That the patient is informed:

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

1 **treating the patient;**

2 **“(B) Of each potential risk associated with receiving the treatment**
3 **known to the consulting physician;**

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

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

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

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

14 **“(G) That expanded access to treating the patient’s terminal disease**
15 **may be provided pursuant to 21 C.F.R. 312.300 to 312.320 and may be**
16 **an option for the patient, and that:**

17 **“(i) A patient who receives treatment pursuant to 21 C.F.R. 312.300**
18 **to 312.320 is not required to pay the costs of administering the treat-**
19 **ment or the costs of, or the costs associated with, manufacturing the**
20 **investigational product as described in section 3 (1)(b) of this 2015 Act;**
21 **and**

22 **“(ii) An insurer is not relieved of reimbursing costs as described in**
23 **section 12 of this 2015 Act for treatment provided pursuant to 21 C.F.R.**
24 **312.300 to 312.320.**

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

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 consulting physician a patient is suf-
5 fering from a psychiatric or psychological disorder or depression
6 causing impaired judgment, the consulting physician shall refer the
7 patient for counseling. Treatment may not be provided as described in
8 section 3 of this 2015 Act until the person performing the counseling
9 determines that the patient is not suffering from a psychiatric or
10 psychological disorder or depression causing impaired judgment.

11 “SECTION 5. Upon receiving an offer for treatment as described in
12 section 3 of this 2015 Act, a patient who has a terminal disease and
13 who is qualified may elect to receive that treatment by signing and
14 dating a form attesting to the election in the presence of two wit-
15 nesses. A form attesting to an election must include:

16 “(1) The attending physician’s diagnosis for the patient;

17 “(2) The attending physician’s prognosis for the patient;

18 “(3) A statement that the investigational product to be used in
19 treating the patient is not approved by the United States Food and
20 Drug Administration;

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

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

26 “(a) Any health care practitioner who participates in administering
27 the treatment, to whom a health care practitioner who participates in
28 administering the treatment refers the patient or with whom a health
29 care practitioner who participates in administering the treatment
30 consults;

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

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

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

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

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

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

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

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

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

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

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

25 **“SECTION 7. A waiver of liability required by section 5 (5) of this**
26 **2015 Act must be written in plain and simple language.**

27 **“SECTION 8. (1) Except as provided in subsection (3) of this section,**
28 **a health care practitioner who participates in administering a treat-**
29 **ment as described in section 3 of this 2015 Act, or a health care facility**
30 **or professional organization or association involved in the adminis-**

1 tration of the treatment, is not subject to civil or criminal liability for
2 acts or omissions of acts related to administering the treatment if the
3 administration of the treatment complies with sections 1 to 14 of this
4 2015 Act.

5 “(2) Except as provided in subsection (3) of this section, a man-
6 ufacturer or distributor of an investigational product used to treat a
7 patient pursuant to section 3 of this 2015 Act is not subject to civil or
8 criminal liability for acts or omissions of acts related to the adminis-
9 tration of the investigational product.

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

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

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

22 “SECTION 10. A health care facility or health care practitioner may
23 prohibit another health care practitioner from participating in ad-
24 ministering a treatment as described in section 3 of this 2015 Act at
25 the health care facility or on premises owned or controlled by the
26 prohibiting health care practitioner.

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

29 “(1) A licensing board, health care facility, health care practitioner
30 or professional organization or association may impose upon the vio-

1 **lating health care practitioner any form of discipline referred to in**
2 **section 9 of this 2015 Act that the licensing board, health care facility,**
3 **health care practitioner or professional organization or association**
4 **otherwise may legally impose; and**

5 **“(2) The health care facility or prohibiting health care practitioner**
6 **may:**

7 **“(a) Terminate any lease or other property contract entered into**
8 **with the violating health care practitioner and subject the violating**
9 **health care practitioner to any other nonmonetary remedies provided**
10 **by such a contract; or**

11 **“(b) Terminate any contract for the provision of services entered**
12 **into with the violating health care practitioner and subject the vio-**
13 **lating health care practitioner to any other nonmonetary remedies**
14 **provided by such a contract.**

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

17 **“(1) Associated with undergoing a treatment as described in section**
18 **3 of this 2015 Act; or**

19 **“(2) Demonstrated by clear and convincing evidence to be associated**
20 **with an adverse effect that is a result of undergoing a treatment as**
21 **described in section 3 of this 2015 Act.**

22 **“SECTION 13. Eligibility for hospice care must be determined on**
23 **the basis of a patient’s overall prognosis and care or treatment goals**
24 **as determined by the patient’s attending physician and may not be**
25 **determined on the basis of whether a patient is undergoing or has**
26 **undergone a treatment as described in section 3 of this 2015 Act.**

27 **“SECTION 14. (1) The Oregon Health Authority shall annually re-**
28 **view a sample of records maintained pursuant to sections 1 to 14 of**
29 **this 2015 Act.**

30 **“(2) An attending physician who makes a referral under section 2**

1 of this 2015 Act, a health care practitioner who administers treatment
2 as described in section 3 of this 2015 Act and a consulting physician
3 who provides written confirmation as described in section 4 of this 2015
4 Act must file with the authority a record, in a form and manner pre-
5 scribed by the authority, of the findings of the attending physician,
6 health care practitioner or consulting physician.

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

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

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

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

13 “(d) The demographics of the patients to whom the treatment is
14 administered.

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

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

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

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

30