

**PROPOSED AMENDMENTS TO
A-ENGROSSED HOUSE BILL 3345**

1 On page 1 of the printed A-engrossed bill, line 12, after “facility” insert
2 “, as those terms are defined in ORS 442.015”.

3 In line 13, delete “or”.

4 In line 14, delete the period and insert “; or

5 “(D) A clinical laboratory, as defined in ORS 438.010, that is:

6 “(i) Licensed under ORS 438.010 to 438.510; and

7 “(ii) Owned or controlled by, or under common ownership with, a hospital
8 described in subparagraph (A) of this paragraph.”.

9 In line 16, delete the colon.

10 In line 17, delete “(a)”.

11 In line 18, delete the semicolon and insert a period.

12 Delete lines 19 through 26.

13 On page 2, delete lines 1 through 14.

14 In line 15, delete “(3)” and insert “(2)”.

15 In line 18, delete “(4)” and insert “(3)”.

16 In line 20, delete “(5)” and insert “(4)”.

17 In line 22, delete “(6)” and insert “(5)”.

18 On page 3, line 5, before “The” insert “(1)”.

19 In line 6, delete “(1)” insert “(a)” and delete “and”.

20 In line 7, delete “(2)” and insert “(b)”.

21 In line 9, delete the period and insert “; and

22 “(c) In lieu of conducting inspections authorized under paragraph (b) of

1 this subsection, accept accreditation from an accrediting body approved by
2 the authority.

3 “(2) To be approved under subsection (1)(c) of this section, an accrediting
4 body must:

5 “(a) Require a nontransplant anatomical research recovery organization
6 to document processes related to the recovery, handling and distribution of
7 anatomical material and submit to the organization that documentation.

8 “(b) Require a nontransplant anatomical research recovery organization
9 to keep and maintain all records related to the recovery or distribution of
10 anatomical material for at least 10 years.

11 “(c) Conduct, or have a designee conduct, regular on-site compliance in-
12 spections of a nontransplant anatomical research recovery organization’s
13 records, processes and materials relating to:

14 “(A) Donor intake;

15 “(B) Acquisition, preparation, labeling, packaging, storage and distrib-
16 ution of anatomical material; and

17 “(C) Any inspection of a facility owned or operated by the nontransplant
18 anatomical research recovery organization.

19 “(d) Meet any other requirement adopted by the authority by rule.”.

20 In line 21, after “licensed” delete the rest of the line.

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