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LEGISLATIVE ACTION

Senate	.	House
Comm: RCS	.	
02/27/2020	.	
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The Committee on Rules (Hutson) recommended the following:

Senate Amendment (with title amendment)

Delete everything after the enacting clause
and insert:

Section 1. Section 381.06017, Florida Statutes, is created to read:

381.06017 Nonembryonic stem cell banks; collecting, manufacturing, storing, dispensing, and using adult human nonembryonic stem cells and HCT/Ps.—

(1) DEFINITIONS.—As used in this section, the term:

(a) "Adult human nonembryonic stem cells" means cells that

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are derived from adult human nonembryonic HCT/Ps through enzymatic digestion, mechanical disruption, or similar processing. The term includes only drugs, devices, or biological products that are approved by the United States Food and Drug Administration and are regulated by the FD&C Act, s. 351 of the PHS Act, or part I of chapter 499.

(b) "Agency" means the Agency for Health Care Administration.

(c) "Allogenic use" means the collection of human cells or tissue from one person and the implantation, transplantation, infusion, or transfer of those human cells or tissue into another person.

(d) "Autologous use" means the implantation, transplantation, infusion, or transfer of human cells or tissue back into the individual from which they were collected.

(e) "Dispense" has the same meaning as in s. 465.003(6).

(f) "Establishment" means a place of business which is at one general physical location and may extend to one or more contiguous suites, units, floors, or buildings operated and controlled exclusively by entities under common operation and control. The term includes multiple buildings with an intervening thoroughfare if the buildings are under common exclusive ownership, operation, and control. For purposes of permitting, each suite, unit, floor, or building must be identified in the most recent permit application.

(g) "FD&C Act" means the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. ss. 301 et seq.; 52 Stat. 1040 et seq.

(h) "HCT/Ps" means human cells, tissues, or cellular or tissue-based products that are intended for implantation,

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transplantation, infusion, or transfer into a human recipient. The term does not include any of the following:

1. Vascularized human organs for transplantation.

2. Whole blood, blood components, blood derivative products, or platelet-rich plasma that are exempt under 21 C.F.R 607.65.

3. Human secretions, including milk, collagen, and cell factors, but not semen.

4. Minimally manipulated bone marrow that is for homologous use only and that is not combined with any other article except water, crystalloids, or sterilizing, preserving, or storage agents.

5. Ancillary products used in the manufacture of nonembryonic adult human allogenic or autologous HCT/Ps.

6. Cells, tissue, or organs derived from animals.

7. In vitro diagnostic products.

8. Blood vessels recovered with an organ for transplantation.

(i) "Homologous use" means the repair, reconstruction, or supplementation of a recipient's cells or tissues with adult human nonembryonic stem cells or adult human nonembryonic HCT/Ps that perform the same basic function or functions in the recipient as in the donor.

(j) "Manufacture" means the preparing, deriving, compounding, propagation, processing, producing, or fabricating of any drug, device, or cosmetic.

(k) "Minimally manipulated" means:

1.a. For structural tissues, processing that does not alter the original characteristics of the tissue which relate to the

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tissue's utility for reconstruction, repair, or replacement; or

b. For cells or nonstructural tissues, processing that does not alter the relevant biological characteristics of the cells or tissues.

2. The washing, rinsing, cleaning, sizing, shaping, or concentrating of adult human nonembryonic HCT/Ps which does not alter the relevant characteristics or basic functions of the tissue or cell.

(l) "Nonembryonic stem cell bank" means a publicly or privately owned establishment that operates its own laboratories, retains control over all aspects of processing and storage, is managed by a single entity, and performs any of the following activities in the course of its business:

1. Engages in the manufacture, use, implantation, transplantation, infusion, dispensing, transfer, or storage of adult human allogenic and autologous nonembryonic stem cells.

2. Accepts, receives, carries, or delivers human allogenic and autologous nonembryonic stem cells, drugs, or products that are approved by United States Food and Drug Administration and regulated as drugs, devices, or biological products by the FD&C Act, s. 251 of the PHS Act, or part I of chapter 499.

3. Recovers, collects, screens, and tests, in the facility, adult human autologous nonembryonic HCT/Ps from a specific patient for implantation, transplantation, infusion, or transfer back into the same patient during a single surgery within the facility.

4. Provides patient-specific health care services using adult human autologous nonembryonic HCT/Ps in the facility during a single procedure.


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5. Advertises adult human nonembryonic stem cell services or adult human autologous nonembryonic HCT/P services, including, but not limited to, the collection, manufacture implantation, transplantation, infusion, transfer, storage, dispensing, use, or purported use of United States Food and Drug Administration-approved adult human autologous nonembryonic stem cells or adult human autologous nonembryonic HCT/Ps that are intended to diagnose, cure, mitigate, treat, provide therapy for, or prevent an injury or a disease.

6. Performs any procedure that is intended to:

a. Collect or store adult human autologous nonembryonic HCT/Ps for autonomous homologous use; or

b. Diagnose, cure, mitigate, treat, provide therapy for, or prevent an injury or a disease through the use or purported use of adult human autologous nonembryonic HCT/Ps.

7. Compounds patient-specific adult human autologous nonembryonic HCT/Ps into a drug product by combining or mixing the patient-specific adult human nonembryonic HCT/Ps, at the prescriptive direction of a licensed physician authorized within the scope of his or her license to prescribe and administer adult human autologous nonembryonic HTC/Ps with one or more drugs or products to create a patient-specific drug or product.

8. Dispenses adult human autologous nonembryonic stem cells or HTC/Ps to any of the following for a specific patient pursuant to a valid order from a licensed physician authorized within the scope of his or her license to prescribe and administer adult human autologous nonembryonic HTC/Ps:

a. The specific patient’s physician with privileges to practice at the nonembryonic stem cell bank.

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b. For office use, the specific patient's physician's office or a health care facility or treatment setting where the physician has privileges to administer adult human autologous nonembryonic HTC/Ps.

(l) "Office use" includes the provision and administration of any United States Food and Drug Administration-approved adult human nonembryonic stem cell drug, compounded drug, or compounded product regulated as a drug, device, or any biological product under the FD&C Act, the PHS Act, 42 U.S.C. 262, s. 351, or part I of chapter 499, to a patient's physician in the physician's office or in a health care facility or treatment setting, including a hospital, an ambulatory surgical center, or a health care clinic licensed under chapter 395 or chapter 400. The term also includes the patient-specific dispensing, provision, or administration of the patient's adult human autologous nonembryonic HTC/Ps.

(m) "PHS Act" means the Public Health and Safety Act, 42 U.S.C. ss. 262 et seq., and applicable regulations, including 21 C.F.R. part 1271.

(n) "Physician" means a person who is licensed to practice medicine under chapter 458 or osteopathic medicine under chapter 459.

(2) DUTIES AND REGISTRATION.—

(a) Establishments that manufacture adult human nonembryonic HTC/Ps are regulated by s. 361 of the PHS Act and part I of chapter 499. Such establishments must register with and submit a list of all HCT/Ps manufactured to the Food and Drug Administration and obtain a permit from the Department of Business and Professional Regulation if the HCT/P manufactured

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is:

1. Minimally manipulated;

2. Intended only for homologous use;

3. Manufactured through a process that does not involve the combination of the cells or tissue with another article, except water, crystalloids, or a sterilizing, preserving, or storing agent; and

4. For an adult human nonembryonic HCT/P, either:

a. Does not have a systemic effect and is not dependent upon the metabolic activity of living cells for their primary function; or

b. Has a systemic effect or is dependent upon the metabolic activity of living cells for its primary function and is for autologous use or for allogenic use in a first-degree or second-degree blood relative.

(b) Establishments that manufacture adult human nonembryonic HCT/Ps that do not meet the criteria described in paragraph (a) are exempt from the registration and listing requirements of s. 361 of the PHS Act, but must obtain a permit from and submit a list of all HTC/Ps manufactured to the Department of Business and Professional Regulation if the establishment:

1. Uses the adult human nonembryonic HTC/Ps for nonmedicinal scientific purposes; or

2. Removes human adult nonembryonic HCT/Ps from a patient and implants the same HCT/Ps into the same patient during the same surgical procedure with only minimal manipulation of the HCT/Ps which does not alter the original relevant biological characteristics of the cells or tissues.

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(c) A nonembryonic stem cell bank that manufactures adult human nonembryonic HCT/Ps may not more than minimally manipulate, through enzymatic digestion, mechanical disruption, or similar processing, any adult human nonembryonic stem cell or HCT/P to alter the HCT/P's original structural characteristics or relevant biological characteristics or to isolate differentiated cells from undifferentiated cells that have lost their original structural function, so that the undifferentiated cells can be differentiated into a specialized cell type, unless the nonembryonic stem cell bank has first registered the HCT/P with the United States Food and Drug Administration and the Department of Business and Professional Regulation as a drug, device, or biological product manufacturer and complies with all applicable regulations under the FD&C Act, s. 351 of the PHS Act, 21 C.F.R. parts 1-1299, and part I of chapter 499.

(d) A nonembryonic stem cell bank that advertises, collects, stores, manufactures, dispenses, compounds, uses, or purports to use adult human nonembryonic stem cells or adult human autologous nonembryonic HCT/Ps is deemed a clinic as defined in s. 400.9905 and must comply with all of the following requirements:

1. Adhere to the applicable current good manufacturing practices for the collecting, removing, manufacturing, processing, using, compounding, and implanting of adult human nonembryonic stem cells or products containing adult human nonembryonic stem cells pursuant to the FD&C Act, the PHS Act, 21 C.F.R., parts 1270-1271, and part I of chapter 499.

2. Adhere to the applicable current good manufacturing practices for the collecting, removing, manufacturing,

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processing, using, compounding, and implanting of adult human autologous nonembryonic HCT/Ps so that it does not alter the relevant tissue or cellular characteristics or basic functions.

3. Obtain a health care clinic license from the agency pursuant to s. 400.991 and part II of chapter 408 and register each establishment separately, unless:

a. The clinic is a facility licensed under chapter 395; or

b. The clinic is affiliated with an accredited medical school that provides training to medical students, residents, or fellows.

4. Have a physician medical director who is responsible for the establishment's compliance with all requirements related to licensure, operation of a nonembryonic stem cell bank, and current good manufacturing practices under this section, part X of chapter 400, and the FD&C Act, the PHS Act, 21 C.F.R. parts 1-1299, and part I of chapter 499.

5. Notify the agency, in writing, on a form approved by the agency, within 10 days after termination of a physician medical director and notify the agency within 10 days after such termination of the identity of the physician medical director who has assumed responsibility for that nonembryonic stem cell bank. Failure to have a physician medical director practicing at the location of the licensed nonembryonic stem cell bank is the basis for a summary suspension of the nonembryonic stem cell bank's license pursuant to s. 120.60(6) or s. 400.607.

6. Require a physician medical director with a full, active, and unencumbered license to actively practice at the nonembryonic stem cell bank location for which he or she has assumed responsibility.


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7. Maintain commercial and professional liability insurance in an amount not less than $250,000 per claim.

8. Operate each establishment using the same name as the one used to obtain the health care clinic license from the agency. All invoices, packing slips, and other business records must list the same name.

9. Obtain a pharmacy permit for each person and establishment before dispensing, offering office use of, or compounding adult human nonembryonic stem cells with any other drug, compound, or product.

(3) DISPENSING OF DRUGS OR COMPOUNDED DRUGS OR PRODUCTS.—

(a) A pharmacist at a nonembryonic stem cell bank that is also permitted as a pharmacy under chapter 465 may dispense for office use only any of the following to a stem cell bank within this state:

1. Adult human nonembryonic stem cells.

2. A compounded drug containing adult human nonembryonic stem cells.

3. A compounded product containing adult human nonembryonic stem cells.

(b) Adult human nonembryonic stem cells, compounded drugs containing adult human nonembryonic stem cells, or products containing adult human nonembryonic stem cells may not be sold or dispensed by any person or establishment other than the adult human nonembryonic stem cell bank or a pharmacist at the nonembryonic stem cell bank that dispenses or receives the adult human nonembryonic stem cells or the compounded drug or product containing adult human nonembryonic stem cells, except that:

1. A physician who requests the dispensing of adult human

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nonembryonic stem cells, a compounded drug, or a compounded product from the manufacturing nonembryonic stem cell bank may administer such items to his or her patient if the physician is authorized within the scope of his or her license to prescribe and administer adult human nonembryonic stem cells; or

2. A pharmacist, a pharmacy, or an establishment that receives or carries adult human nonembryonic stem cells, a compounded drug, or a compounded product that was manufactured by a nonembryonic stem cell bank may sell or dispense such items to a physician who is authorized within the scope of his or her license to prescribe and administer adult human nonembryonic stem cells to patients.

(4) HEALTH CARE PRACTITIONER RESPONSIBILITIES.—

(a) A physician, an advanced practice registered nurse licensed under chapter 464, or a physician assistant licensed under chapter 458 or chapter 459 may not practice in a nonembryonic stem cell bank that is not licensed with the agency as required by the rules adopted pursuant to s. 400.9925. The license of a health care practitioner who violates this paragraph is subject to disciplinary action by the appropriate regulatory board.

(b) In the performance of any procedure collecting, storing, using, or purporting to use adult human nonembryonic stem cells or products containing adult human nonembryonic stem cells, a health care practitioner must adhere to the applicable current good manufacturing practices for the collecting, removing, manufacturing, processing, using, compounding, and implanting of stem cells or products containing stem cells pursuant to the FD&C Act, 21 C.F.R., parts 1270-1271, the PHS

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Act, and part I of chapter 499.

(5) RULEMAKING.—The agency, in consultation with the Department of Health and the Department of Business and Professional Regulation, shall adopt rules necessary to administer the licensure, inspection, and regulation of nonembryonic stem cell banks, including, but not limited to, rules regarding all of the following which must be consistent with the best practices specified in the FD&C Act, 21 C.F.R. parts 1270-1271, the PHS Act, and part I of chapter 499:

(a) Advertising.

(b) Nonembryonic stem cell bank procedures and protocols for the collecting, removing, manufacturing, storing, dispensing, and using of adult human nonembryonic stem cells, other drugs containing adult human nonembryonic stem cells, and products containing adult human nonembryonic stem cells, in accordance with applicable current best practices.

(c) Adverse incident reporting.

(d) Informed consent.

(e) Recordkeeping, record retention, and availability of records for inspection.

Section 2. This act shall take effect July 1, 2020.

================ T I T L E A M E N D M E N T ================

And the title is amended as follows:

Delete everything before the enacting clause and insert:

A bill to be entitled

An act relating to nonembryonic stem cell banks; creating s. 381.06017, F.S.; defining terms; providing

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registration requirements for certain establishments; prohibiting a nonembryonic stem cell bank from more than minimally manipulating adult human nonembryonic stem cells or HCT/Ps under certain circumstances; providing that a nonembryonic stem cell bank that performs certain functions is deemed a clinic; requiring such nonembryonic stem cell banks to comply with specified requirements; prohibiting an entity other than certain nonembryonic stem cell banks and pharmacists from dispensing certain compounded drugs or products, with exceptions; prohibiting certain health care practitioners from practicing in a nonembryonic stem cell bank that is not licensed with the agency; providing for disciplinary action; requiring health care practitioners to adhere to specified regulations in the performance of certain procedures; requiring the Agency, in consultation with the Department of Health and the Department of Business and Professional Regulation, to adopt specified rules; providing an effective date.