From House Calendar, 5/1/14, Pg. 2908-2910

H. 350
An act relating to the posting of medical unprofessional conduct decisions and to investigators of alleged unprofessional conduct

The Senate proposes to the House to amend the bill as follows:

First: In Sec. 2, 26 V.S.A. § 1368, in subdivision (a)(4)(B), at the end of the subdivision following ―within five business days of the expiration of the appeal period‖, by inserting or within five business days of the request of the licensee, whichever is later.

Second: By adding a new section to be numbered Sec. 5a to read as follows:

Sec. 5a. 18 V.S.A. § 4631a is amended to read:

§ 4631a. EXPENDITURES BY MANUFACTURERS OF PRESCRIBED PRODUCTS

(a) As used in this section:

(1) ―Allowable expenditures‖ means:

* * *

(H) Sponsorship of an educational program offered by a medical device manufacturer at a national or regional professional society meeting at which programs accredited by the Accreditation Council for Continuing Medical Education, or a comparable professional accrediting entity, are also offered, provided:

(i) no payment is made directly to a health care professional or pharmacist; and

(ii) the funding is used solely for bona fide educational purposes, except that the manufacturer may provide meals and other food for program participants.

(I) Other reasonable fees, payments, subsidies, or other economic benefits provided by a manufacturer of prescribed products at fair market value.

* * *

(7)(C) ―Regularly practices‖ means to practice at least periodically under contract with, as an employee of, or as the owner of, a medical practice, health care facility, nursing home, hospital, or university located in Vermont.

* * *

(12) ―Prescribed product‖ means a drug ~~or device~~ as defined in section 201 of the federal Food, Drug and Cosmetic Act, 21 U.S.C. § 321, a compound drug or drugs, a medical device as defined in this subsection, a biological product as defined in section 351 of the Public Health Service Act, 42 U.S.C. § 262, for human use, or a combination product as defined in 21 C.F.R. § 3.2(e), but shall not include prescription eyeglasses, prescription sunglasses, or other prescription eyewear.

* * *

(15) ―Medical device‖ means an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent, or other similar or related article, including any component, part, or accessory, that is:

(A) recognized in the official National Formulary or the United States Pharmacopeia, or any supplement to them;

(B) intended for use in the diagnosis of disease or other conditions, or in the cure, mitigation, treatment, or prevention of disease, in humans or other animals; or

(C) intended to affect the structure or any function of the body of humans or other animals, and which does not achieve its primary intended purposes through chemical action within or on such body and which is not dependent upon being metabolized for the achievement of its primary intended purposes.

Third: By striking out Sec. 6 (Effective Dates) in its entirety and inserting in lieu thereof a new Sec. 6 to read as follows:

Sec. 6. EFFECTIVE DATES

This act shall take effect on passage, except:

(1) Secs. 1 (amending 26 V.S.A. § 1318), 3 (amending 26 V.S.A. § 1351), and 5a (amending 18 V.S.A. § 4631a) shall take effect on July 1, 2014; and

(2) Sec. 2 (amending 26 V.S.A. § 1368) shall take effect on July 1, 2015.

(For text see House Journal January 23, 2014)