

S.31

An act relating to informed health care financial decision making

The House proposes to the Senate to amend the bill by striking all after the enacting clause and inserting in lieu thereof the following:

Sec. 1. 18 V.S.A. chapter 42 is amended to read:

CHAPTER 42. BILL OF RIGHTS FOR HOSPITAL PATIENTS AND  
PATIENT ACCESS TO INFORMATION

Subchapter 1. Bill of Rights for Hospital Patients

§ 1851. DEFINITIONS

As used in this ~~chapter~~ subchapter:

(1) "Hospital" means a ~~general~~ hospital required to be licensed under ~~18 V.S.A. chapter 43~~ of this title.

(2) "Patient" means a person admitted to a hospital on an inpatient basis.

§ 1852. PATIENTS' BILL OF RIGHTS; ADOPTION

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(12) The patient has the right to receive an itemized, detailed, and understandable explanation of charges regardless of the source of payment and to be provided with information about financial assistance and billing and collections practices.

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Subchapter 2. Access to Information

§ 1854. PUBLIC ACCESS TO INFORMATION

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§ 1855. AMBULATORY SURGICAL PATIENTS; EXPLANATION OF  
CHARGES

(a) As used in this section:

(1) “Ambulatory surgical center” has the same meaning as in section  
9432 of this title.

(2) “Hospital” means a hospital required to be licensed under chapter 43  
of this title.

(b) A patient receiving outpatient surgical services or an outpatient  
procedure at an ambulatory surgical center or hospital shall receive an  
itemized, detailed, and understandable explanation of charges regardless of the  
source of payment and shall be provided with information about the  
ambulatory surgical center’s or hospital’s financial assistance and billing and  
collections practices.

Sec. 2. 18 V.S.A. § 9375(b) is amended to read:

(b) The Board shall have the following duties:

\* \* \*

(14) Collect and review data from each psychiatric hospital licensed  
pursuant to chapter 43 of this title, which may include data regarding a

psychiatric hospital's scope of services, volume, utilization, discharges, payer mix, quality, coordination with other aspects of the health care system, and financial condition. The Board's processes shall be appropriate to psychiatric hospitals' scale and their role in Vermont's health care system, and the Board shall consider ways in which psychiatric hospitals can be integrated into systemwide payment and delivery system reform.

Sec. 3. 18 V.S.A. § 9351 is amended to read:

§ 9351. HEALTH INFORMATION TECHNOLOGY PLAN

(a)(1) The Department of Vermont Health Access, in consultation with the Department's Health Information Exchange Steering Committee, shall be responsible for the overall coordination of Vermont's statewide Health Information Technology Plan. The Plan shall be revised annually and updated comprehensively every five years to provide a strategic vision for clinical health information technology.

(2) The Department shall submit the proposed Plan to the Green Mountain Care Board annually on or before November 1. The Green Mountain Care Board shall approve, reject, or request modifications to the Plan within 45 days following its submission; if the Board has taken no action after 45 days, the Plan shall be deemed to have been approved.

(3)(A) The Department, in consultation with the Steering Committee, shall administer the Plan, ~~which shall~~.

(B) The Plan shall include the implementation of an integrated electronic health information infrastructure for the sharing of electronic health information among health care facilities, health care professionals, public and private payers, and patients. The Plan shall provide for each patient's electronic health information to be accessible to health care facilities, health care professionals, and public and private payers to the extent permitted under federal law unless the patient has affirmatively elected not to have the patient's electronic health information shared in that manner.

(C) The Plan shall include standards and protocols designed to promote patient education, patient privacy, physician best practices, electronic connectivity to health care data, access to advance care planning documents, and, overall, a more efficient and less costly means of delivering quality health care in Vermont.

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Sec. 4. VERMONT HEALTH INFORMATION EXCHANGE; OPT-OUT  
CONSENT POLICY; IMPLEMENTATION

(a) The Department of Vermont Health Access, in consultation with its Health Information Exchange Steering Committee, shall administer a robust stakeholder process to develop an implementation strategy for the consent policy for the sharing of patient health information through the Vermont

Health Information Exchange (VHIE), as revised pursuant to Sec. 3 of this act.

The implementation strategy shall:

(1) include substantial opportunities for public input;

(2) focus on the creation of patient education mechanisms and processes

that:

(A) combine new information on the consent policy with existing patient education obligations, such as disclosure requirements under the Health Insurance Portability and Accountability Act of 1996 (HIPAA);

(B) aim to address diverse needs, abilities, and learning styles with respect to information delivery;

(C) clearly explain:

(i) the purpose of the VHIE;

(ii) the way in which health information is currently collected;

(iii) how and with whom health information may be shared using the VHIE;

(iv) the purposes for which health information may be shared using the VHIE;

(v) how to opt out of having health information shared using the VHIE; and

(vi) how patients can change their participation status in the future; and

(D) enable patients to fully understand their rights regarding the sharing of their health information and provide them with ways to find answers to associated questions, including providing contact information for the Office of the Health Care Advocate;

(3) identify the mechanisms by which Vermonters will be able to easily opt out of having their health information shared through the VHIE and a timeline identifying when each mechanism will be available, which shall begin in advance of the July 1, 2020 change to the consent policy;

(4) include plans for developing or supplementing consent management processes at the VHIE to reflect the needs of patients and providers;

(5) include multisector communication strategies to inform each Vermonter about the VHIE, the consent policy, and their ability to opt out of having their health information shared through the VHIE; and

(6) identify a methodology for evaluating the extent to which the public outreach regarding the VHIE, consent policy, and opt-out processes has been successful.

(b)(1) The Department of Vermont Health Access shall provide updates on the stakeholder engagement process and the consent policy implementation strategy to the House Committee on Health Care, the Senate Committee on Health and Welfare, the Health Reform Oversight Committee, and the Green Mountain Care Board on or before August 1 and November 1, 2019.

(2) The Department of Vermont Health Access shall provide a final report on the outcomes of the stakeholder engagement process and the consent policy implementation strategy to the House Committee on Health Care, the Senate Committee on Health and Welfare, and the Green Mountain Care Board on or before January 15, 2020.

Sec. 5. EFFECTIVE DATES

(a) Secs. 1 (18 V.S.A. chapter 42) and 2 (18 V.S.A. § 9375(b) shall take effect on July 1, 2019.

(b) Sec. 3 (18 V.S.A. § 9351) shall take effect on July 1, 2020.

(c) Sec. 4 (Vermont Health Information Exchange; opt-out consent policy; implementation) and this section shall take effect on passage.

and that after passage the title of the bill be amended to read: “An act relating to informed health care financial decision making and the consent policy for the Vermont Health Information Exchange”