

1 TO THE HOUSE OF REPRESENTATIVES:

2 The Committee on Health Care to which was referred Senate Bill No. 31  
3 entitled “An act relating to informed health care financial decision making”  
4 respectfully reports that it has considered the same and recommends that the  
5 House propose to the Senate that the bill be amended by striking out all after  
6 the enacting clause and inserting in lieu thereof the following:

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

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

10 Subchapter 1. Bill of Rights for Hospital Patients

11 § 1851. DEFINITIONS

12 As used in this ~~chapter~~ subchapter:

13 (1) “Hospital” means a ~~general~~ hospital required to be licensed under  
14 ~~18 V.S.A. chapter 43~~ of this title.

15 (2) “Patient” means a person admitted to a hospital on an inpatient basis.

16 § 1852. PATIENTS’ BILL OF RIGHTS; ADOPTION

17 \* \* \*

18 (12) The patient has the right to receive an itemized, detailed, and  
19 understandable explanation of charges regardless of the source of payment and  
20 to be provided with information about financial assistance and billing and  
21 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:

\* \* \*

1           (14) Collect and review data from each psychiatric hospital licensed  
2           pursuant to chapter 43 of this title, which may include data regarding a  
3           psychiatric hospital’s scope of services, volume, utilization, discharges, payer  
4           mix, quality, coordination with other aspects of the health care system, and  
5           financial condition. The Board’s processes shall be appropriate to psychiatric  
6           hospitals’ scale and their role in Vermont’s health care system, and the Board  
7           shall consider ways in which psychiatric hospitals can be integrated into  
8           systemwide payment and delivery system reform.

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

10          § 9351. HEALTH INFORMATION TECHNOLOGY PLAN

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

17           (2) The Department shall submit the proposed Plan to the Green  
18           Mountain Care Board annually on or before November 1. The Green  
19           Mountain Care Board shall approve, reject, or request modifications to the Plan  
20           within 45 days following its submission; if the Board has taken no action after  
21           45 days, the Plan shall be deemed to have been approved.



1 policy for the sharing of patient health information through the Vermont  
2 Health Information Exchange (VHIE), as revised pursuant to Sec. 3 of this act.

3 The implementation strategy shall:

4 (1) include substantial opportunities for public input;

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

6 that:

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

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

12 (C) clearly explain:

13 (i) the purpose of the VHIE;

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

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

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

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

- 1                    (vi) how patients can change their participation status in the  
2 future; and
- 3                    (D) enable patients to fully understand their rights regarding the  
4 sharing of their health information and provide them with ways to find answers  
5 to associated questions, including providing contact information for the Office  
6 of the Health Care Advocate;
- 7                    (3) identify the mechanisms by which Vermonters will be able to easily  
8 opt out of having their health information shared through the VHIE and a  
9 timeline identifying when each mechanism will be available, which shall begin  
10 in advance of the July 1, 2020 change to the consent policy;
- 11                    (4) include plans for developing or supplementing consent management  
12 processes at the VHIE to reflect the needs of patients and providers;
- 13                    (5) include multisector communication strategies to inform each  
14 Vermonters about the VHIE, the consent policy, and their ability to opt out of  
15 having their health information shared through the VHIE; and
- 16                    (6) identify a methodology for evaluating the extent to which the public  
17 outreach regarding the VHIE, consent policy, and opt-out processes has been  
18 successful.
- 19                    (b)(1) The Department of Vermont Health Access shall provide updates on  
20 the stakeholder engagement process and the consent policy implementation  
21 strategy to the House Committee on Health Care, the Senate Committee on

1 Health and Welfare, the Health Reform Oversight Committee, and the Green  
2 Mountain Care Board on or before August 1 and November 1, 2019.

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

8 Sec. 5. EFFECTIVE DATES

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

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

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

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

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18 (Committee vote: \_\_\_\_\_)

19 \_\_\_\_\_

20 Representative \_\_\_\_\_

21 FOR THE COMMITTEE