

1 Committee of Conference Report on HB 1466, relative to the off-label use of prescription drugs and
2 relative to pharmacy prescriptions.

3
4 Recommendation:

5 That the House recede from its position of nonconcurrency with the Senate amendment, and
6 concur with the Senate amendment, and

7 That the Senate and House adopt the following new amendment to the bill as amended by the
8 Senate, and pass the bill as so amended:

9
10 Amend the bill by replacing sections 1-3 with the following:

11
12 1 New Paragraph; Physicians and Surgeons; Disciplinary Action; Off-Label Use of Prescription
13 Drug. Amend RSA 329:17 by inserting after paragraph VI-a the following new paragraph:

14 VI-b. The state of New Hampshire confirms its strong support for shared decision making
15 between healthcare professionals and their patients. A licensee may lawfully prescribe an FDA
16 approved drug product for an off-label indication and be held to the same standard of care as when
17 prescribing for on-label indication when:

18 (a) Off-label use of the drug product for this indication has longstanding common use;

19 (b) There is medical evidence to support this use and no known evidence
20 contraindicating such use, including but not limited to peer reviewed studies and practice guidelines
21 from relevant medical societies; or

22 (c) The licensee has provided and the patient, or if the patient is a minor, the patient's
23 parent or guardian, has signed an informed consent form that includes the known potential benefits,
24 known potential risks, alternative treatment options, expected prognosis without treatment, and a
25 disclosure that a prescription is for an off-label indication. The signed informed consent form shall
26 remain part of the patient's medical record.

27 2 New Paragraph; Nurse Practice Act; Advanced Practice Registered Nurse; Disciplinary Action;
28 Off-label Use of Prescription Drug. Amend RSA 326-B:37 by inserting after paragraph III the
29 following new paragraph:

30 III-a. The state of New Hampshire confirms its strong support for shared decision making
31 between healthcare professionals and their patients. A licensee may lawfully prescribe an FDA
32 approved drug product for an off-label indication and be held to the same standard of care as when
33 prescribing for on-label indication when:

34 (a) Off-label use of the drug product for this indication has longstanding common use;

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1 (b) There is medical evidence to support this use and no known evidence
2 contraindicating such use, including but not limited to peer reviewed studies and practice guidelines
3 from relevant medical societies; or

4 (c) The licensee has provided and the patient, or if the patient is a minor, the patient's
5 parent or guardian, has signed an informed consent form that includes the known potential benefits,
6 known potential risks, alternative treatment options, expected prognosis without treatment, and a
7 disclosure that a prescription is for an off-label indication. The signed informed consent form shall
8 remain part of the patient's medical record.

9 3 New Section; Physician Assistant; Disciplinary Action; Off-label Use of Prescription Drug.
10 Amend RSA 328-D by inserting after section 6 the following new section:

11 328-D:6-a Off-label Use of Prescription Drugs; When Permitted. The state of New Hampshire
12 confirms its strong support for shared decision making between healthcare professionals and their
13 patients. A licensee may lawfully prescribe an FDA approved drug product for an off-label indication
14 and be held to the same standard of care as when prescribing for on-label indication when:

15 I. Off-label use of the drug product for this indication has longstanding common use;

16 II. There is medical evidence to support this use and no known evidence contraindicating
17 such use, including but not limited to peer reviewed studies and practice guidelines from relevant
18 medical societies; or

19 III. The licensee has provided and the patient, or if the patient is a minor, the patient's
20 parent or guardian, has signed an informed consent form that includes the known potential benefits,
21 known potential risks, alternative treatment options, expected prognosis without treatment, and a
22 disclosure that a prescription is for an off-label indication. The signed informed consent form shall
23 remain part of the patient's medical record.

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The signatures below attest to the authenticity of this Report on HB 1466, relative to the off-label use of prescription drugs and relative to pharmacy prescriptions.

Conferees on the Part of the Senate

Conferees on the Part of the House

Sen. Bradley, Dist. 3

Rep. Layon, Rock. 6

Sen. Avard, Dist. 12

Rep. Merchant, Sull. 4

Sen. Sherman, Dist. 24

Rep. Kofalt, Hills. 4

Rep. B. King, Hills. 23