



State of Wisconsin
2017 - 2018 LEGISLATURE

LRBa0066/1
MED:jld

**ASSEMBLY AMENDMENT 2,
TO ASSEMBLY SUBSTITUTE AMENDMENT 1,
TO ASSEMBLY BILL 69**

March 1, 2017 - Offered by Representatives SUBECK, KOLSTE, C. TAYLOR and
ZAMARRIPA.

- 1 At the locations indicated, amend the substitute amendment as follows:
- 2 **1.** Page 3, line 17: after "MANUFACTURERS." insert "(a)".
- 3 **2.** Page 3, line 19: after "patient." insert:
- 4 "(b)".
- 5 **3.** Page 3, line 23: after that line insert:
- 6 "(c) During the time an eligible patient is using an investigational drug, device,
- 7 or biological product as provided in this section, the manufacturer shall notify the
- 8 treating physician and the eligible patient of any clinically significant finding of a
- 9 known or anticipated risk, side effect, or reported patient discomfort that is likely
- 10 related to the investigational drug, device, or biological product that becomes known

1 to the manufacturer after the eligible patient begins using the investigational drug,
2 device, or biological product.”.

3 (END)