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| PREVAILED | Roll Call No. _____ |
| FAILED | Ayes _____ |
| WITHDRAWN | Noes _____ |
| RULED OUT OF ORDER | |

# HOUSE MOTION ____

MR. SPEAKER:

I move that Engrossed Senate Bill 3 be amended to read as follows:

Page 3, between lines 10 and 11, begin a new paragraph and insert:

"**(e) When issuing a standing order, prescription, or protocol to administer or dispense an immunization that is recommended by the federal Centers for Disease Control and Prevention Advisory Committee on Immunization Practices or the federal Food and Drug Administration, the state health commissioner or the commissioner's designated public health authority shall do the following:**

**(1) Commission an independent third party review by a licensed medical professional for the immunization of all adverse event reports from the state that were submitted to the national Vaccine Adverse Event Reporting System (VAERS). A review by a third party concerning submitted adverse events by individuals in the state must:**

**(A) be conducted not later than thirty (30) days from submission by an individual of an adverse event to VAERS;**

**(B) include a follow up phone call to each individual who has submitted an adverse event at least quarterly for twelve (12) months unless the individual refuses further contact, in order to ensure appropriate follow up concerning the immunization; and**

**(C) the provision of notice concerning the Countermeasures Injury Compensation Program (CICP) or the Vaccine Injury Compensation Program (VICP).**

**The third party reviewer shall submit a quarterly summary of the aggregate data from the review that does not include personal identifying information to the state department and the general assembly in an electronic format under IC 5-14-6. A person conducting a third party review under this subdivision may not release personal identifying information concerning an individual who filed an adverse event report unless the person obtains written consent from the individual who made the adverse event report. A person conducting a third party review under this subdivision, upon the request of an individual who has submitted an adverse event report concerning the immunization being reviewed, may include written testimony and supplemental documentation as part of the report from the individual who reported the adverse event.**

**(2) Issue an official advisory from the state department to health care providers performing immunizations under the standing order, prescription, or protocol described in this section that includes the following requirements of a health care provider:**

**(A) The standing order, prescription, or protocol to administer an immunization.**

**(B) The health care provider's legal responsibility concerning reporting adverse events to the national Vaccine Adverse Event Reporting System (VAERS)**.

**(C) A list of possible side effects of the immunization that have been reported for the immunization or similar immunizations.**

**(D) Information concerning the Countermeasures Injury Compensation Program (CICP) and the Vaccine Injury Compensation Program (VICP) and the importance of educating patients.**

**(E) A statement that a third party review may be performed on the immunization being received and to cooperate with the review if contacted.**

**(f) A health care provider that administers an immunization under a standing order, prescription, or protocol described in this section must do the following:**

**(1) Train staff on the following:**

**(A) The national Vaccine Adverse Event Reporting System (VAERS)**.

**(B) The Countermeasures Injury Compensation Program (CICP).**

**(C) The Vaccine Injury Compensation Program (VICP).**

**(2) Post information concerning the programs described in subdivision (1) in the area in which the health care provider is providing the immunizations.**

**(3) Educate patients verbally concerning the programs described in subdivision (1) before providing the immunization.**

**(4) Have immunization manufacturer inserts that may be provided to the patient upon the patient's request.**".

Page 3, line 11, delete "(e)" and insert "**(g)**".

Page 3, line 11, after "(b)," insert "**and subject to subsection (e),**".

(Reference is to ESB 3 as printed February 17, 2022.)

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Representative Nisly