1 ENGROSSED HOUSE BILL NO. 3791 By: Marti, Davis and Virgin of 2 the House 3 and 4 Standridge of the Senate 5 6 An Act relating to pharmacy; defining terms; providing the substitution of an interchangeable biological product for a prescribed biological 7 product under certain conditions; requiring electronic notice of substitution; providing that 8 dispensing pharmacist shall not be required to show 9 certain proof of access; providing exceptions; directing State Board of Pharmacy to maintain link of 10 all interchangeable biological products; providing for approved brand and generic substitutions; providing for codification; and providing an 11 effective date. 12 1.3 14 15 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 16 SECTION 1. A new section of law to be codified NEW LAW 17 in the Oklahoma Statutes as Section 353.18B of Title 59, unless 18 there is created a duplication in numbering, reads as follows: 19 A. As used in this section: 20 "Biological product" has the same meaning given to that term 21 in 42 U.S.C., Section 262; and 22 "Interchangeable biological product" means a biological 23 product that the United States Food and Drug Administration (USFDA):

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- a. has licensed and determined to meet the standards for interchangeability pursuant to 42 U.S.C., Section 262(k)(4) of the Internal Revenue Code, or
  - b. has determined is therapeutically equivalent as set forth in the latest edition of or supplement to the USFDA's Approved Drug Products with Therapeutic Equivalence Evaluations, commonly known as the Orange Book.
- B. A pharmacist may substitute an interchangeable biological product for a prescribed biological product only if all of the following conditions are met:
- 1. The substituted product has been determined by the USFDA to be interchangeable, as defined in subsection A of this section, with the prescribed biological product;
  - 2. The prescribing physician has permitted substitution; and
  - 3. The pharmacy informs the patient of the substitution.
- C. Within five (5) business days following the dispensing of a biological product, the dispensing pharmacist or the pharmacist's designee shall make an entry of the specific product provided to the patient, including the name of the product and the manufacturer. The communication shall be conveyed by making an entry that can be electronically accessed by the prescriber through:
  - 1. An interoperable electronic medical records system;
  - 2. An electronic prescribing technology;

- 3. A pharmacy benefit management system; or
  - 4. A pharmacy record.

- D. The dispensing pharmacist or a prescriber shall not be:
- 1. Required to show proof that the prescriber has access to the record in any type of payment audit conducted by a payer or pharmacy benefit manager; or
- 2. Subject to disciplinary action or civil penalties for failure to ensure that the record is accessible or for failure to access the record.
- E. Entry into an electronic records system as described in subsection C of this section is presumed to provide notice to the prescriber. Otherwise, the pharmacist shall communicate the biological product dispensed to the prescriber using facsimile, telephone, electronic transmission or other prevailing means, except that communication shall not be required when:
- 1. There is no USFDA-approved interchangeable biological product for the product prescribed; or
- 2. A refill prescription is not changed from the product dispensed on the prior filling of the prescription.
- F. The State Board of Pharmacy shall maintain a link on its
  Internet website to the current list of all biological products
  determined by the USFDA to be interchangeable with a specific
  biological product.

1	G. Nothing in this section shall preclude existing approved
2	brand and generic substitutions.
3	SECTION 2. This act shall become effective November 1, 2020.
4	Passed the House of Representatives the 3rd day of March, 2020.
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6	Presiding Officer of the House
7	of Representatives
8	Passed the Senate the day of, 2020.
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