1	ENGROSSED SENATE
2	BILL NO. 1341 By: McCortney of the Senate
	and
3	McEntire of the House
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6	An Act relating to pharmacy; providing definitions; allowing a pharmacist to substitute certain
7	interchangeable product for certain prescribed product if certain conditions are met; requiring a
8	pharmacist or designee to make entry of certain product provided within certain time frame; providing
9	for method of certain communication; providing for notice to certain prescriber; providing exemption for
10	certain communication; excluding dispensing pharmacist or prescriber from certain requirement and
11	certain penalties; directing the State Board of
12	Pharmacy to maintain certain link on its website; providing certain construction; providing for
13	codification; and providing an effective date.
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15	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:
16	SECTION 1. NEW LAW A new section of law to be codified
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. For the purposes of this section:
20	1. "Biological product" has the same meaning given to that term
21	in 42 U.S.C., Section 262; and
22	2. "Interchangeable biological product" means a biological
23	product that the U.S. Food and Drug Administration (FDA):

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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), or
 - b. has determined is therapeutically equivalent as set forth in the latest edition of or supplement to the United States Food and Drug Administration's (FDA) Approved Drug Products with Therapeutic Equivalence Evaluations.
- B. A pharmacist may substitute an interchangeable biological product for a prescribed biological product only if all of the following conditions in this subsection are met:
- 1. The substituted product has been determined by FDA 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.

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- D. 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 where:
 - 1. There is no FDA-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.
 - E. 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.
- 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 FDA to be interchangeable with a specific
 biological product.

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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 Senate the 3rd day of March, 2020.
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6	Drociding Officer of the Consta
7	Presiding Officer of the Senate
8	Passed the House of Representatives the day of,
9	2020.
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