1		AN.	ACT relating to the practice of pharmacy.	
2	Be it enacted by the General Assembly of the Commonwealth of Kentucky:			
3		<b>→</b> Se	ection 1. KRS 315.010 is amended to read as follows:	
4	As u	sed in	this chapter, unless the context requires otherwise:	
5	(1)	"Adı	minister" means the direct application of a drug to a patient or research subject	
6		by ir	njection, inhalation, or ingestion, whether topically or by any other means;	
7	(2)	"Administrative activities of a pharmacy" means the following functions performed		
8		by a	pharmacy adhering to all local, state, and federal patient privacy laws:	
9		(a)	Investigating and researching a patient's insurance benefits and updating the	
10			patient profile regarding insurance coverage;	
11		(b)	Billing and collections activities, including:	
12			1. Contacting patients for copayments and coinsurance payments; and	
13			2. Communicating with insurance companies;	
14		(c)	Performing patient financial assistance activities and updating patient records	
15			accordingly;	
16		(d)	Opening faxes and accessing electronic prescriptions for the purposes of	
17			setting up patient demographic and insurance profiles, excluding height,	
18			weight, and allergy information, so long as the activity does not involve the	
19			entering of a prescription order into the dispensing or medication management	
20			system;	
21		(e)	Initiating insurance prior authorizations for submission to the licensed	
22			pharmacy, including communications with the prescribing physician to	
23			collect, record, and transmit information to insurance companies, so long as	
24			the activity does not include the authorization or receipt of new or refill	
25			prescription orders;	
26		(f)	Answering and transferring telephone calls, whether or not such calls require	

accessing a patient record, so long as the call does not involve the

1			interpretation, evaluation, or implementation of a drug order; and
2		(g)	Communicating with patients via telephone or electronically regarding refill
3			reminders, so long as the communication does not involve the interpretation,
4			evaluation, or implementation of a drug order and a pharmacist is readily
5			available for patient consultation;
6	(3)	"Ass	sociation" means the Kentucky Pharmacists Association;
7	(4)	"Bo	ard" means the Kentucky Board of Pharmacy;
8	(5)	"Co	laborative care agreement" means a written agreement between a pharmacist or
9		phar	macists and a practitioner or practitioners that outlines a plan of cooperative
0		man	agement of patients' drug-related health care needs where:
1		(a)	Patients' drug-related health care needs fall within the practitioner's or
12			practitioners' statutory scope of practice;
13		(b)	Patients are referred by the practitioner or practitioners to the pharmacist or
4			pharmacists; and
5		(c)	The agreement:
6			1. Identifies the practitioner or practitioners and the pharmacist or
17			pharmacists who are parties to the agreement;
8			2. Specifies the drug-related regimen to be provided, and how drug therapy
9			is to be monitored; and
20			3. Stipulates the conditions for initiating, continuing, or discontinuing drug
21			therapy and conditions which warrant modifications to dose, dosage
22			regimen, dosage form, or route of administration;
23	(6)	"Co	mpound" or "compounding" means the preparation or labeling of a drug
24		purs	uant to or in anticipation of a valid prescription drug order, including but not
25		limi	ted to packaging, intravenous admixture, or manual combination of drug
26		ingr	edients. "Compounding," as used in this chapter, shall not preclude simple
27		reco	nstitution, mixing, or modification of drug products prior to administration by

1		nonp	pharmacists;
2	(7)	"Cor	nfidential information" means information which is accessed or maintained by a
3		phar	macist in a patient's record, or communicated to a patient as part of patient
4		coun	seling, whether it is preserved on paper, microfilm, magnetic media, electronic
5		medi	ia, or any other form;
6	(8)	"Cor	ntinuing education unit" means ten (10) contact hours of board approved
7		conti	inuing pharmacy education. A "contact hour" means fifty (50) continuous
8		minu	ites without a break period;
9	(9)	"Dis	pense" or "dispensing" means to deliver one (1) or more doses of a prescription
10		drug	in a suitable container, appropriately labeled for subsequent administration to
11		or us	se by a patient or other individual entitled to receive the prescription drug;
12	(10)	"Dru	g" means any of the following:
13		(a)	Articles recognized as drugs or drug products in any official compendium or
14			supplement thereto;
15		(b)	Articles, other than food, intended to affect the structure or function of the
16			body of man or other animals;
17		(c)	Articles, including radioactive substances, intended for use in the diagnosis,
18			cure, mitigation, treatment or prevention of disease in man or other animals;
19			or
20		(d)	Articles intended for use as a component of any articles specified in
21			paragraphs (a) to (c) of this subsection;
22	(11)	"Dru	ng regimen review" means retrospective, concurrent, and prospective review by
23		a ph	armacist of a patient's drug-related history, including but not limited to the

- 25 (a) Evaluation of prescription drug orders and patient records for:
- 26 1. Known allergies;

following areas:

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27 2. Rational therapy contraindications;

1			3. Appropriate dose and route of administration;
2			4. Appropriate directions for use; or
3			5. Duplicative therapies;
4		(b)	Evaluation of prescription drug orders and patient records for drug-drug,
5			drug-food, drug-disease, and drug-clinical laboratory interactions;
6		(c)	Evaluation of prescription drug orders and patient records for adverse drug
7			reactions; or
8		(d)	Evaluation of prescription drug orders and patient records for proper
9			utilization and optimal therapeutic outcomes;
10	(12)	"Imr	mediate supervision" means under the physical and visual supervision of a
11		phar	macist;
12	(13)	"Ma	nufacturer" or "virtual manufacturer" of a product means:
13		(a)	A person that holds an application approved under 21 U.S.C. sec. 355 or a
14			license issued under 42 U.S.C. sec. 262 for such product, or if such product is
15			not the subject of an approved application or license, the person who
16			manufactured the product;
17		(b)	A co-licensed partner of the person described in paragraph (a) of this
18			subsection that obtains the product directly from a person described in this
19			paragraph or paragraph (a) of this subsection;
20		(c)	An affiliate of a person described in paragraph (a) or (b) of this subsection
21			who receives the product directly from a person described in this paragraph or
22			in paragraph (a) or (b) of this subsection; or
23		(d)	Any person, except a pharmacist compounding in the normal course of
24			professional practice;
25	(14)	"Me	dical order" means a lawful order of a specifically identified practitioner for a
26		spec	ifically identified patient for the patient's health care needs. "Medical order"
27		may	or may not include a prescription drug order;

1	(15)	"Nor	apprescription drugs" means nonnarcotic medicines or drugs which may be sold	
2		with	out a prescription and are prepackaged and labeled for use by the consumer in	
3		accordance with the requirements of the statutes and regulations of this state and the		
4		fede	ral government;	
5	(16)	"Out	sourcing facility" means a facility at one (1) geographic location or address	
6		that:		
7		(a)	Is engaged in the compounding of human sterile drugs without a patient-	
8			specific prescription;	
9		(b)	Has registered as an outsourcing facility with the secretary of the United	
10			States Department of Health and Human Services, Food and Drug	
11			Administration; and	
12		(c)	Complies with all applicable state and federal requirements;	
13	(17)	"Pha	rmacist" means a natural person licensed by this state to engage in the practice	
14		of th	e profession of pharmacy;	
15	(18)	"Pha	rmacist intern" means a natural person who is:	
16		(a)	Currently certified by the board to engage in the practice of pharmacy under	
17			the direction of a licensed pharmacist and who satisfactorily progresses	
18			toward meeting the requirements for licensure as a pharmacist;	
19		(b)	A graduate of an approved college or school of pharmacy or a graduate who	
20			has established educational equivalency by obtaining a Foreign Pharmacy	
21			Graduate Examination Committee (FPGEC) certificate, who is currently	
22			licensed by the board for the purpose of obtaining practical experience as a	
23			requirement for licensure as a pharmacist;	
24		(c)	A qualified applicant awaiting examination for licensure as a pharmacist or	
25			the results of an examination for licensure as a pharmacist; or	
26		(d)	An individual participating in a residency or fellowship program approved by	
27			the board for internship credit;	

- 1 (19) "Pharmacy" means every place where:
- 2 (a) Drugs are dispensed under the direction of a pharmacist;
- 3 (b) Prescription drug orders are compounded under the direction of a pharmacist;

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- 5 (c) A registered pharmacist maintains patient records and other information for 6 the purpose of engaging in the practice of pharmacy, whether or not 7 prescription drug orders are being dispensed;
- 8 (20) "Pharmacy-related primary care" means the pharmacists' activities in patient 9 education, health promotion, and assistance in the selection and use of over-the-10 counter drugs and appliances for the treatment of common diseases and injuries, as 11 well as those other activities falling within their statutory scope of practice;
  - (21) "Pharmacy technician" means a natural person who works under the immediate supervision, or general supervision if otherwise provided for by statute or administrative regulation, of a pharmacist for the purpose of assisting a pharmacist with the practice of pharmacy;
  - (22) "Practice of pharmacy" means interpretation, evaluation, and implementation of medical orders and prescription drug orders; responsibility for dispensing prescription drug orders, including radioactive substances; participation in drug and drug-related device selection; administration of medications or biologics in the course of dispensing or maintaining a prescription drug order; the administration of adult immunizations pursuant to prescriber-approved protocols; the administration of immunizations to individuals *three*[nine] (3[9]) to seventeen (17) years of age pursuant to prescriber-approved protocols with the consent of a parent or guardian; the administration of immunizations to a child as defined in KRS 214.032, pursuant to protocols as authorized by KRS 315.500; drug evaluation, utilization, or regimen review; maintenance of patient pharmacy records; and provision of patient counseling and those professional acts, professional decisions, or professional

1		services necessary to maintain and manage all areas of a patient's pharmacy-related		
2		care, including pharmacy-related primary care as defined in this section;		
3	(23)	"Practitioner" has the same meaning given in KRS 217.015(35);		
4	(24)	"Prescription drug" means a drug which:		
5		(a) Under federal law is required to be labeled with either of the following		
6		statements:		
7		1. "Caution: Federal law prohibits dispensing without prescription";		
8		2. "Caution: Federal law restricts this drug to use by, or on the order of, a		
9		licensed veterinarian";		
0		3. "Rx Only"; or		
1		4. "Rx"; or		
12		(b) Is required by any applicable federal or state law or administrative regulation		
13		to be dispensed only pursuant to a prescription drug order or is restricted to		
4		use by practitioners;		
5	(25)	"Prescription drug order" means an original or new order from a practitioner for		
6		drugs, drug-related devices or treatment for a human or animal, including orders		
17		issued through collaborative care agreements or protocols authorized by the board.		
8		Lawful prescriptions result from a valid practitioner-patient relationship, are		
9		intended to address a legitimate medical need, and fall within the prescribing		
20		practitioner's scope of professional practice;		
21	(26)	"Society" means the Kentucky Society of Health-Systems Pharmacists;		
22	(27)	"Supervision" means the presence of a pharmacist on the premises to which a		
23		pharmacy permit is issued, who is responsible, in whole or in part, for the		
24		professional activities occurring in the pharmacy; and		
25	(28)	"Wholesaler" means any person who legally buys drugs for resale or distribution to		

persons other than patients or consumers.

→ Section 2. KRS 315.205 is amended to read as follows:

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1 Upon the request of an individual or his or her parent or guardian, a pharmacist who

- 2 administers an immunization to an individual who is <u>three</u>[nine] (<u>3[9]</u>) to seventeen (17)
- 3 years of age, as authorized in KRS 315.010[(22)], shall provide notification of the
- 4 immunization to the individual's primary care provider.