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1 AN ACT relating to blood donation.

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## 2 Be it enacted by the General Assembly of the Commonwealth of Kentucky:

- 3 → Section 1. KRS 214.452 is amended to read as follows:
- 4 The following policies shall apply to blood establishments and to donors of blood:
- 5 All blood establishments within the Commonwealth shall be licensed by the United (1)6 States Food and Drug Administration and remain in compliance with all applicable 7 federal regulations. The Cabinet for Health and Family Services shall, under 8 administrative regulations promulgated pursuant to KRS Chapter 13A, establish 9 fees necessary to cover the cost of and adhere to a schedule for regular inspection, 10 by the Office of the Inspector General of the Cabinet for Health and Family 11 Services, of all blood establishments within the Commonwealth to ascertain 12 whether each blood establishment is licensed and in compliance with KRS 214.450 13 to 214.464 and KRS 214.468. The Office of the Inspector General shall commence 14 its inspection program of blood establishments no later than September 1, 1994.
- (2) All blood establishments shall test blood for the human immunodeficiency virus 16 and for any known causative agent for any blood-borne communicable disease, using tests approved and required, for purposes of blood donation, by the United States Food and Drug Administration.
- 19 (3)It shall be the duty of the administrator of any blood establishment which collects 20 blood for the purpose of distributing to another health service, health facility, or 21 health-care provider the blood for transfusion to:
  - Secure donor consent and a signed written risk factor history and donor (a) consent form for each potential paid or volunteer donor for the purpose of determining if the potential donor:
- 25 Is at high risk for infection with the human immunodeficiency virus; f, or <u>1.</u> 26 }
- 27 Has tested confirmatory positive for infection with the human <u>2.</u>

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I			immunodeficiency virus; [ or ]
2			3. Has acquired immune deficiency syndrome; [ or ]
3			4. Has tested confirmatory positive for infection with any causative agent
4			for acquired immune deficiency syndrome recognized by the United
5			States Centers for Disease Control; [or]
6			5. Has a blood-borne communicable disease; or
7			6. Has received a COVID-19 vaccine;
8		(b)	Provide a means for a potential donor to self-elect not to donate blood;
9		(c)	Refuse donation or sale of blood by persons at high risk for infection with the
10			human immunodeficiency virus, or who have been medically diagnosed as
11			having acquired immune deficiency syndrome, or who have tested
12			confirmatory positive for infection with the human immunodeficiency virus,
13			or who have a blood-borne communicable disease; and
14		(d)	Post a sign in the blood establishment which is visible to all potential donors
15			and which states: "Persons with acquired immune deficiency syndrome
16			(AIDS), or who have tested confirmatory positive for infection with the
17			human immunodeficiency virus (HIV), or who have a blood-borne
18			communicable disease or who have one (1) or more risk factors for the human
19			immunodeficiency virus as determined by the United States Centers for
20			Disease Control, are prohibited by law from donating or selling blood.
21			Persons violating the law are guilty of a Class D felony. ASK STAFF OF
22			THIS BLOOD ESTABLISHMENT."
23	(4)	The	provisions of this section shall not be construed to impose requirements which
24		are i	n conflict with donor eligibility requirements set out in United States Food and
25		Dru	g Administration or American Association of Blood Banks standards.
26		<b>→</b> S	ection 2. KRS 214.458 is amended to read as follows:
27	(1)	Eacl	n unit of blood collected by a blood establishment for transfusion shall be

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1	affixed	with	the	United	States	Food	and	Drug .	Administ	ration	requir	ed lal	oel
2	and[wh	<del>ich in</del>	clude	es] a do	onor id	entifica	tion	number	through	which	the fe	ollowi	ng

- 3 information can be obtained:
- 4 (a) *The* date the blood was collected;
- 5 (b) <u>The</u> name of <u>the</u> blood establishment;
- 6 (c) A nonidentifying code representing the name of the blood donor;
- 7 (d) A blood establishment serial number for the blood;
- 8 (e) The date of laboratory testing of the blood;
- 9 (f) The name of the person and laboratory testing the blood;
- 10 (g) The laboratory test results; *and*

## 11 (h) Notice that the blood donor signed a statement that he or she had received a

## 12 <u>COVID-19 vaccine</u>.

- 13 Each unit of blood received by a blood establishment or health facility within the (2) 14 Commonwealth from an out-of-state blood establishment shall contain a label in 15 accordance with the provisions of subsection (1) of this section and the blood establishment or health facility shall either test the blood in accordance with the 16 17 requirements for blood establishments within the Commonwealth under the 18 provisions of KRS 214.452(2) or may accept documented evidence of the test 19 results as are required under subsection (1) of this section for blood collected within 20 the Commonwealth.
- 21 (3) Each laboratory testing blood for transfusion shall maintain for ten (10) years from 22 the date of testing, and each blood establishment shall maintain for ten (10) years 23 from the date of collection, a list containing the information set forth in subsection 24 (1) of this section.
- No blood may be transfused into any patient in any health facility or health service or by any health care provider unless the unit of blood has affixed to it the label as required under this section and the blood has tested negative for the human

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1	11	mmunodeficiency virus or any causative agent of AIDS <u>and</u> [ <del>, or]</del> any blood-borne
2	c	communicable disease as provided under KRS 214.452.
3	(5) <b>E</b>	Except in an emergency, before blood is transfused into any patient, a health
4	£	acility, health service, or a health care provider shall inform the patient if the
5	<u>b</u>	lood donor signed a statement that he or she had received a COVID-19 vaccine.
6	<u>(6)</u> V	When a unit of blood is transfused, a label containing the donor identification
7	n	number required under this section shall be removed from the unit and affixed to
8	tl	he patient's medical chart or the blood donor identification number for the unit of
9	b	plood shall be recorded in the patient's medical chart.
10	<u>(7)</u> [(5)	Any unit of blood not containing the label required under this section shall be
11	d	lestroyed by the health facility, health service, or health care provider.
12	<u>(8)</u> [(6)	Any unit of blood testing confirmatory positive for an agent of a blood-borne
13	c	communicable disease and in the possession of a health facility, health service, or
14	h	nealth care provider may be donated to educational or scientific research
15	iı	nstitutions for the purpose of scientific research only and not for transfusion.