^{116TH CONGRESS} 2D SESSION **S. 4547**

To improve the integrity and safety of horseracing by requiring uniform safety and performance standards, including a horseracing anti-doping and medication control program and a racetrack safety program to be developed and enforced by an independent Horseracing Integrity and Safety Authority, and for other purposes.

IN THE SENATE OF THE UNITED STATES

SEPTEMBER 9, 2020

Mr. MCCONNELL (for himself, Mrs. GILLIBRAND, Ms. MCSALLY, and Mrs. FEINSTEIN) introduced the following bill; which was read twice and referred to the Committee on Commerce, Science, and Transportation

A BILL

- To improve the integrity and safety of horseracing by requiring uniform safety and performance standards, including a horseracing anti-doping and medication control program and a racetrack safety program to be developed and enforced by an independent Horseracing Integrity and Safety Authority, and for other purposes.
 - 1 Be it enacted by the Senate and House of Representa-
 - 2 tives of the United States of America in Congress assembled,

3 SECTION 1. SHORT TITLE.

4 This Act may be cited as the "Horseracing Integrity

5 and Safety Act of 2020".

1 SEC. 2. DEFINITIONS.

2 In this Act:

3 (1) AUTHORITY.—The term "Authority" means 4 the Horseracing Integrity and Safety Authority des-5 ignated by section 3(a). COMMISSION.—The term "Commission" 6 (2)7 means the Federal Trade Commission. COVERED HORSE.—The term "covered 8 (3)horse" means any Thoroughbred horse, or any other 9 10 horse made subject to this Act by election of the ap-11 plicable State racing commission or the breed gov-12 erning organization for such horse under section 13 5(k), during the period—

14 (A) beginning on the date of the horse's
15 first timed and reported workout at a racetrack
16 that participates in covered horseraces or at a
17 training facility; and

(B) ending on the date on which the Authority receives written notice that the horse
has been retired.

(4) COVERED HORSERACE.—The term "covered
horserace" means any horserace involving covered
horses that has a substantial relation to interstate
commerce, including any Thoroughbred horserace
that is the subject of interstate off-track or advance
deposit wagers.

1 (5) COVERED PERSONS.—The term "covered 2 persons" means all trainers, owners and breeders, 3 jockeys, racetracks, veterinarians, persons (legal and 4 natural) licensed by a State racing commission and 5 the agents, assigns, and employees of such persons 6 and other horse support personnel who are engaged 7 in the care, training, or racing of covered horses.

8 (6) EQUINE CONSTITUENCIES.—The term 9 "equine constituencies" means, collectively, owners 10 and breeders, trainers, racetracks, veterinarians, 11 State racing commissions, and jockeys who are en-12 gaged in the care, training, or racing of covered 13 horses.

14 (7) Equine industry representative.—The term "equine industry representative" means an or-15 ganization regularly and significantly engaged in the 16 17 equine industry, including organizations that rep-18 resent the interests of, and whose membership con-19 sists of, owners and breeders, trainers, racetracks, 20 veterinarians, State racing commissions, and jock-21 eys.

(8) HORSERACING ANTI-DOPING AND MEDICATION CONTROL PROGRAM.—The term "horseracing anti-doping and medication control program" means

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1	the anti-doping and medication program established
2	under section $6(a)$.
3	(9) Immediate family member.—The term
4	"immediate family member" shall include a spouse,
5	domestic partner, mother, father, aunt, uncle, sib-
6	ling, or child.
7	(10) INTERSTATE OFF-TRACK WAGER.—The
8	term "interstate off-track wager" has the meaning
9	given such term in section 3 of the Interstate Horse-
10	racing Act of 1978 (15 U.S.C. 3002).
11	(11) Jockey.—The term "jockey" means a
12	rider or driver of a covered horse in covered
13	horseraces.
14	(12) Owners and breeders.—The term
15	"owners and breeders" means those persons who ei-
16	ther hold ownership interests in covered horses or
17	who are in the business of breeding covered horses.
18	(13) Program effective date.—The term
19	"program effective date" means the earlier of—
20	(A) January 1 of the second year after the
21	date of the enactment of this Act; or
22	(B) the date that is 540 days after such
23	date of enactment.

1	(14) RACETRACK.—The term "racetrack"
2	means an organization licensed by a State racing
3	commission to conduct covered horseraces.
4	(15) RACETRACK SAFETY PROGRAM.—The term
5	"racetrack safety program" means the program es-
6	tablished under section 7(a).
7	(16) STAKES RACE.—The term "stakes race"
8	means any race so designated by the racetrack at
9	which such race is run, including, without limitation,
10	the races comprising the Breeders' Cup World
11	Championships and the races designated as graded
12	stakes by the American Graded Stakes Committee of
13	the Thoroughbred Owners and Breeders Association.
14	(17) STATE RACING COMMISSION.—The term
15	"State racing commission" means an entity des-
16	ignated by State law or regulation that has jurisdic-
17	tion over the conduct of horseracing within the ap-
18	plicable State.
19	(18) TRAINER.—The term "trainer" means an
20	individual engaged in the training of covered horses.
21	(19) TRAINING FACILITY.—The term "training
22	facility" means a location that is not a racetrack li-
23	censed by a State racing commission that operates
24	primarily to house covered horses and conduct offi-
25	cial timed workouts.

(20) VETERINARIAN.—The term "veterinarian"
 means a licensed veterinarian who provides veteri nary services to covered horses.

4 (21) WORKOUT.—The term "workout" means a 5 timed running of a horse over a predetermined dis-6 tance not associated with a race or its first quali-7 fying race, if such race is made subject to this Act 8 by election under section 5(k) of the horse's breed 9 governing organization or the applicable State racing 10 commission.

11 SEC. 3. RECOGNITION OF THE HORSERACING INTEGRITY 12 AND SAFETY AUTHORITY.

(a) IN GENERAL.—The private, independent, selfregulatory, nonprofit corporation, to be known as the
"Horseracing Integrity and Safety Authority", is recognized for purposes of developing and implementing a
horseracing anti-doping and medication control program
and a racetrack safety program for covered horses, covered persons, and covered horseraces.

20 (b) BOARD OF DIRECTORS.—

(1) MEMBERSHIP.—The Authority shall be governed by a board of directors (in this section referred to as the "Board") comprised of nine members as follows:

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1	(A) INDEPENDENT MEMBERS.—Five mem-
2	bers of the Board shall be independent mem-
3	bers selected from outside the equine industry.
4	(B) INDUSTRY MEMBERS.—
5	(i) IN GENERAL.—Four members of
6	the Board shall be industry members se-
7	lected from among the various equine con-
8	stituencies.
9	(ii) Representation of equine
10	CONSTITUENCIES.—The industry members
11	shall be representative of the various
12	equine constituencies, and shall include not
13	more than one industry member from any
14	one equine constituency.
15	(2) CHAIR.—The chair of the Board shall be an
16	independent member described in paragraph $(1)(A)$.
17	(3) BYLAWS.—The Board of the Authority shall
18	be governed by bylaws for the operation of the Au-
19	thority with respect to—
20	(A) the administrative structure and em-
21	ployees of the Authority;
22	(B) the establishment of standing commit-
23	tees;
24	(C) the procedures for filling vacancies on
25	the Board and the standing committees;

1	(D) term limits for members and termi-
2	nation of membership; and
3	(E) any other matter the Board considers
4	necessary.
5	(c) Standing Committees.—
6	(1) ANTI-DOPING AND MEDICATION CONTROL
7	STANDING COMMITTEE.—
8	(A) IN GENERAL.—The Authority shall es-
9	tablish an anti-doping and medication control
10	standing committee, which shall provide advice
11	and guidance to the Board on the development
12	and maintenance of the horseracing anti-doping
13	and medication control program.
13 14	and medication control program. (B) MEMBERSHIP.—The anti-doping and
14	(B) MEMBERSHIP.—The anti-doping and
14 15	(B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be
14 15 16	(B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be comprised of seven members as follows:
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14 15 16 17 18	 (B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be comprised of seven members as follows: (i) INDEPENDENT MEMBERS.—A majority of the members shall be independent
14 15 16 17 18 19	 (B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be comprised of seven members as follows: (i) INDEPENDENT MEMBERS.—A majority of the members shall be independent members selected from outside the equine
 14 15 16 17 18 19 20 	 (B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be comprised of seven members as follows: (i) INDEPENDENT MEMBERS.—A majority of the members shall be independent members selected from outside the equine industry.
 14 15 16 17 18 19 20 21 	 (B) MEMBERSHIP.—The anti-doping and medication control standing committee shall be comprised of seven members as follows: (i) INDEPENDENT MEMBERS.—A majority of the members shall be independent members selected from outside the equine industry. (ii) INDUSTRY MEMBERS.—A minority

1	than one industry member from any one
2	equine constituency.
3	(iii) QUALIFICATION.—A majority of
4	individuals selected to serve on the anti-
5	doping and medication control standing
6	committee shall have significant, recent ex-
7	perience in anti-doping and medication
8	control rules.
9	(C) CHAIR.—The chair of the anti-doping
10	and medication control standing committee
11	shall be an independent member of the Board
12	described in subsection (b)(1)(A).
13	(2) RACETRACK SAFETY STANDING COM-
14	MITTEE.—
15	(A) IN GENERAL.—The Authority shall es-
16	tablish a racetrack safety standing committee,
17	which shall provide advice and guidance to the
18	Board on the development and maintenance of
19	the racetrack safety program.
20	(B) Membership.—The racetrack safety
21	standing committee shall be comprised of seven
22	members as follows:
23	(i) INDEPENDENT MEMBERS.—A ma-
24	

1	members selected from outside the equine
2	industry.
3	(ii) INDUSTRY MEMBERS.—A minority
4	of the members shall be industry members
5	selected to represent the various equine
6	constituencies.
7	(C) CHAIR.—The chair of the racetrack
8	safety standing committee shall be an industry
9	member of the Board described in subsection
10	(b)(1)(B).
11	(d) Nominating Committee.—
12	(1) Membership.—
13	(A) IN GENERAL.—The nominating com-
14	mittee of the Authority shall be comprised of
15	seven independent members selected from busi-
16	ness, sports, and academia.
17	(B) INITIAL MEMBERSHIP.—The initial
18	nominating committee members shall be set
19	forth in the governing corporate documents of
20	the Authority.
21	(C) VACANCIES.—After the initial com-
22	mittee members are appointed in accordance
23	with subparagraph (B), vacancies shall be filled
24	by the Board pursuant to rules established by
25	the Authority.

1	(2) CHAIR.—The chair of the nominating com-
2	mittee shall be selected by the nominating committee
3	from among the members of the nominating com-
4	mittee.
5	(3) Selection of members of the board
6	AND STANDING COMMITTEES.—
7	(A) INITIAL MEMBERS.—The nominating
8	committee shall select the initial members of
9	the Board and the standing committees de-
10	scribed in subsection (c).
11	(B) SUBSEQUENT MEMBERS.—The nomi-
12	nating committee shall recommend individuals
13	to fill any vacancy on the Board or on such
14	standing committees.
15	(e) Conflicts of Interest.—To avoid conflicts of
16	interest, the following individuals may not be selected as
17	a member of the Board or as an independent member of
18	a nominating or standing committee under this section:
19	(1) An individual who has a financial interest
20	in, or provides goods or services to, covered horses.
21	(2) An official or officer—
22	(A) of an equine industry representative;
23	or

1	(B) who serves in a governance or policy-
2	making capacity for an equine industry rep-
3	resentative.
4	(3) An employee of, or an individual who has a
5	business or commercial relationship with, an indi-
6	vidual described in paragraph (1) or (2).
7	(4) An immediate family member of an indi-
8	vidual described in paragraph (1) or (2).
9	(f) FUNDING.—
10	(1) INITIAL FUNDING.—
11	(A) IN GENERAL.—Initial funding to es-
12	tablish the Authority and underwrite its oper-
13	ations before the program effective date shall be
14	provided by loans obtained by the Authority.
15	(B) BORROWING.—The Authority may bor-
16	row funds toward the funding of its operations.
17	(C) ANNUAL CALCULATION OF AMOUNTS
18	REQUIRED.—
19	(i) IN GENERAL.—Not later than the
20	date that is 90 days before the program ef-
21	fective date, and not later than November
22	1 each year thereafter, the Authority shall
23	determine and provide to each State racing
24	commission the estimated amount required
25	from the State—

1	(I) to fund the State's propor-
2	tionate share of the horseracing anti-
3	doping and medication control pro-
4	gram and the racetrack safety pro-
5	gram for the next calendar year; and
6	(II) to liquidate the State's pro-
7	portionate share of any loan or fund-
8	ing shortfall in the current calendar
9	year and any previous calendar year.
10	(ii) BASIS OF CALCULATION.—The
11	amounts calculated under clause (i) shall—
12	(I) be based on—
13	(aa) the annual budget of
14	the Authority for the following
15	calendar year, as approved by the
16	Board; and
17	(bb) the projected amount of
18	covered racing starts for the year
19	in each State; and
20	(II) take into account other
21	sources of Authority revenue.
22	(iii) Requirements regarding
23	BUDGETS OF AUTHORITY.—

	14
1	(I) INITIAL BUDGET.—The initial
2	budget of the Authority shall require
3	the approval of $\frac{2}{3}$ of the Board.
4	(II) SUBSEQUENT BUDGETS.—
5	Any subsequent budget that exceeds
6	the budget of the preceding calendar
7	year by more than 5 percent shall re-
8	quire the approval of $\frac{2}{3}$ of the Board.
9	(iv) RATE INCREASES.—
10	(I) IN GENERAL.—A proposed in-
11	crease in the amount required under
12	this subparagraph shall be reported to
13	the Commission.
14	(II) NOTICE AND COMMENT.—
15	The Commission shall publish in the
16	Federal Register such a proposed in-
17	crease and provide an opportunity for
18	public comment.
19	(2) Assessment and collection of fees by
20	STATES.—
21	(A) NOTICE OF ELECTION.—Any State
22	racing commission that elects to remit fees pur-
23	suant to this subsection shall notify the Author-
24	ity of such election not later than 60 days be-
25	fore the program effective date.

1	(B) REQUIREMENT TO REMIT FEES.—
2	After a State racing commission makes a notifi-
3	cation under subparagraph (A), the election
4	shall remain in effect and the State racing com-
5	mission shall be required to remit fees pursuant
6	to this subsection according to a schedule estab-
7	lished in rule developed by the Authority and
8	approved by the Commission.
9	(C) WITHDRAWAL OF ELECTION.—A State
10	racing commission may cease remitting fees
11	under this subsection not earlier than one year
12	after notifying the Authority of the intent of
13	the State racing commission to do so.
14	(D) DETERMINATION OF METHODS.—Each
15	State racing commission shall determine, sub-
16	ject to the applicable laws, regulations, and con-
17	tracts of the State, the method by which the
18	requisite amount of fees, such as foal registra-
19	tion fees, sales contributions, starter fees, and
20	track fees, and other fees on covered persons,
21	shall be allocated, assessed, and collected.
22	(3) Assessment and collection of fees by
23	THE AUTHORITY.—
24	(A) CALCULATION.—If a State racing com-
25	mission does not elect to remit fees pursuant to

1	paragraph (2) or withdraws its election under
2	such paragraph, the Authority shall, not less
3	frequently than monthly, calculate the applica-
4	ble fee per racing start multiplied by the num-
5	ber of racing starts in the State during the pre-
6	ceding month.
7	(B) Allocation.—The Authority shall al-
8	locate equitably the amount calculated under
9	subparagraph (A) collected among covered per-
10	sons involved with covered horseraces pursuant
11	to such rules as the Authority may promulgate.
12	(C) Assessment and collection.—
13	(i) IN GENERAL.—The Authority shall
14	assess a fee equal to the allocation made
15	under subparagraph (B) and shall collect
16	such fee according to such rules as the Au-
17	thority may promulgate.
18	(ii) REMITTANCE OF FEES.—Covered
19	persons described in subparagraph (B)
20	shall be required to remit such fees to the
21	Authority.
22	(D) LIMITATION.—A State racing commis-
23	sion that does not elect to remit fees pursuant
24	to paragraph (2) or that withdraws its election
25	under such paragraph shall not impose or col-

1	lect from any person a fee or tax relating to
2	anti-doping and medication control or racetrack
3	safety matters for covered horseraces.
4	(4) FEES AND FINES.—Fees and fines imposed
5	by the Authority shall be allocated toward funding
6	of the Authority and its activities.
7	(5) RULE OF CONSTRUCTION.—Nothing in this
8	Act shall be construed to require—
9	(A) the appropriation of any amount to the
10	Authority; or
11	(B) the Federal Government to guarantee
12	the debts of the Authority.
13	(g) QUORUM.—For all items where Board approval
14	is required, the Authority shall have present a majority
15	of independent members.
16	SEC. 4. FEDERAL TRADE COMMISSION OVERSIGHT.
17	(a) IN GENERAL.—The Authority shall submit to the
18	Commission, in accordance with such rules as the Com-
19	mission may prescribe under section 553 of title 5, United
20	States Code, any proposed rule, or proposed modification
21	to a rule, of the Authority relating to—
22	(1) the bylaws of the Authority;
23	(2) a list of permitted and prohibited medica-

tions, substances, and methods, including allowable

1	limits of permitted medications, substances, and
2	methods;
-	(3) laboratory standards for accreditation and
4	protocols;
5	(4) standards for racing surface quality mainte-
6	nance;
7	(5) racetrack safety standards and protocols;
8	(6) a program for injury and fatality data anal-
9	ysis;
10	(7) a program of research and education on
11	safety, performance, and anti-doping and medication
12	control;
13	(8) a description of safety, performance, and
14	anti-doping and medication control rule violations
15	applicable to covered horses and covered persons;
16	(9) a schedule of civil sanctions for violations;
17	(10) a process or procedures for disciplinary
18	hearings; and
19	(11) a formula or methodology for determining
20	assessments described in section 3(f).
21	(b) Publication and Comment.—
22	(1) IN GENERAL.—The Commission shall—
23	(A) publish in the Federal Register each
24	proposed rule or modification submitted under
25	subsection (a); and

(B) provide an opportunity for public com ment.

3 (2) APPROVAL REQUIRED.—A proposed rule, or 4 a proposed modification to a rule, of the Authority 5 shall not take effect unless the proposed rule or 6 modification has been approved by the Commission. 7 (c) DECISION ON PROPOSED RULE OR MODIFICA-8 TION TO A RULE.— 9 (1) IN GENERAL.—Not later than 60 days after 10 the date on which a proposed rule or modification is 11 published in the Federal Register, the Commission 12 shall approve or disapprove the proposed rule or 13 modification. 14 (2) CONDITIONS.—The Commission shall ap-15 prove a proposed rule or modification if the Commis-16 sion finds that the proposed rule or modification is 17 consistent with— 18 (A) this Act; and 19 (B) applicable rules approved by the Com-20 mission. 21 (3) REVISION OF PROPOSED RULE OR MODI-22 FICATION.— 23 (A) IN GENERAL.—In the case of dis-

approval of a proposed rule or modificationunder this subsection, not later than 30 days

1	after the issuance of the disapproval, the Com-
2	mission shall make recommendations to the Au-
3	thority to modify the proposed rule or modifica-
4	tion.
5	(B) RESUBMISSION.—The Authority may
6	resubmit for approval by the Commission a pro-
7	posed rule or modification that incorporates the
8	modifications recommended under subpara-
9	graph (A).
10	(d) Proposed Standards and Procedures.—
11	(1) IN GENERAL.—The Authority shall submit
12	to the Commission any proposed rule, standard, or
13	procedure developed by the Authority to carry out
14	the horseracing anti-doping and medication control
15	program or the racetrack safety program.
16	(2) NOTICE AND COMMENT.—The Commission
17	shall publish in the Federal Register any such pro-
18	posed rule, standard, or procedure and provide an
19	opportunity for public comment.
20	(e) INTERIM FINAL RULES.—The Commission may
21	adopt an interim final rule, to take effect immediately,
22	under conditions specified in section $553(b)(B)$ of title 5,
23	United States Code, if the Commission finds that such a
24	rule is necessary to protect—

25 (1) the health and safety of covered horses; or

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1	(2) the integrity of covered horseraces and wa-
2	gering on those horseraces.
3	SEC. 5. JURISDICTION OF THE COMMISSION AND THE
4	HORSERACING INTEGRITY AND SAFETY AU-
5	THORITY.
6	(a) IN GENERAL.—Beginning on the program effec-
7	tive date, the Commission, the Authority, and the anti-
8	doping and medication control enforcement agency, each
9	within the scope of their powers and responsibilities under
10	this Act, as limited by subsection (j), shall—
11	(1) implement and enforce the horseracing anti-
12	doping and medication control program and the
13	racetrack safety program;
14	(2) exercise independent and exclusive national
15	authority over—
16	(A) the safety, welfare, and integrity of
17	covered horses, covered persons, and covered
18	horseraces; and
19	(B) all horseracing safety, performance,
20	and anti-doping and medication control matters
21	for covered horses, covered persons, and covered
22	horseraces; and
23	(3) have safety, performance, and anti-doping
24	and medication control authority over covered per-

1	sons similar to such authority of the State racing
2	commissions before the program effective date.
3	(b) PREEMPTION.—The rules of the Authority pro-
4	mulgated in accordance with this Act shall preempt any
5	provision of State law or regulation with respect to mat-
6	ters within the jurisdiction of the Authority under this
7	Act, as limited by subsection (j). Nothing contained in this
8	Act shall be construed to limit the authority of the Com-
9	mission under any other provision of law.
10	(c) DUTIES.—
11	(1) IN GENERAL.—The Authority—
12	(A) shall develop uniform procedures and
13	rules authorizing—
14	(i) access to offices, racetrack facili-
15	ties, other places of business, books,
16	records, and personal property of covered
17	persons that are used in the care, treat-
18	ment, training, and racing of covered
19	horses;
20	(ii) issuance and enforcement of sub-
21	poenas and subpoenas duces tecum; and
22	(iii) other investigatory powers of the
23	nature and scope exercised by State racing
24	commissions before the program effective
25	date; and

1 (B) with respect to an unfair or deceptive 2 act or practice described in section 10, may rec-3 ommend that the Commission commence an en-4 forcement action.

5 (2) APPROVAL OF COMMISSION.—The proce6 dures and rules developed under paragraph (1)(A)
7 shall be subject to approval by the Commission in
8 accordance with section 4.

9 (d) REGISTRATION OF COVERED PERSONS WITH AU-10 THORITY.—

(1) IN GENERAL.—As a condition of participating in covered races and in the care, ownership,
treatment, and training of covered horses, a covered
person shall register with the Authority in accordance with rules promulgated by the Authority and
approved by the Commission in accordance with section 4.

(2) AGREEMENT WITH RESPECT TO AUTHORITY
RULES, STANDARDS, AND PROCEDURES.—Registration under this subsection shall include an agreement by the covered person to be subject to and
comply with the rules, standards, and procedures developed and approved under subsection (c).

24 (3) COOPERATION.—A covered person reg25 istered under this subsection shall, at all times—

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1	(A) cooperate with the Commission, the
2	Authority, the anti-doping and medication con-
3	trol enforcement agency, and any respective
4	designee, during any civil investigation; and
5	(B) respond truthfully and completely to
6	the best of the knowledge of the covered person
7	if questioned by the Commission, the Authority,
8	the anti-doping and medication control enforce-
9	ment agency, or any respective designee.
10	(4) FAILURE TO COMPLY.—Any failure of a
11	covered person to comply with this subsection shall
12	be a violation of section $8(a)(2)(G)$.
13	(e) Enforcement of Programs.—
14	(1) ANTI-DOPING AND MEDICATION CONTROL
15	ENFORCEMENT AGENCY.—
16	(A) AGREEMENT WITH USADA.—The Au-
17	thority shall seek to enter into an agreement
18	with the United States Anti-Doping Agency
19	under which the Agency acts as the anti-doping
20	and medication control enforcement agency
21	under this Act for services consistent with the
22	horseracing anti-doping and medication control
23	program.
24	(B) Agreement with other entity.—If
25	the Authority and the United States Anti-

1 Doping Agency are unable to enter into the 2 agreement described in subparagraph (A), the 3 Authority shall enter into an agreement with an 4 entity that is nationally recognized as being a 5 medication regulation agency equal in qualifica-6 tion to the United States Anti-Doping Agency 7 to act as the anti-doping and medication control 8 enforcement agency under this Act for services 9 consistent with the horseracing anti-doping and 10 medication control program.

11 (C) NEGOTIATIONS.—Any negotiations 12 under this paragraph shall be conducted in 13 good faith and designed to achieve efficient, ef-14 fective best practices for anti-doping and medi-15 cation control and enforcement on commercially 16 reasonable terms.

17 (D)ELEMENTS OF AGREEMENT.—Any 18 agreement under this paragraph shall include a 19 description of the scope of work, performance 20 metrics, reporting obligations, budgets of the 21 United States Anti-Doping Agency while acting 22 as the anti-doping and medication control en-23 forcement agency under this Act, a provision 24 for the revision of the agreement to increase in 25 the scope of work, as provided in subsection (k),

1	and any other matter the Authority considers
2	appropriate.
3	(E) DUTIES AND POWERS OF ENFORCE-
4	MENT AGENCY.—The anti-doping and medica-
5	tion control enforcement agency under an
6	agreement under this paragraph shall—
7	(i) serve as the independent anti-
8	doping and medication control enforcement
9	organization for covered horses, covered
10	persons, and covered horseraces, imple-
11	menting the anti-doping and medication
12	control program on behalf of the Author-
13	ity;
14	(ii) ensure that covered horses and
15	covered persons are deterred from using or
16	administering medications, substances, and
17	methods in violation of the rules estab-
18	lished in accordance with this Act;
19	(iii) implement anti-doping education,
20	research, testing, compliance and adjudica-
21	tion programs designed to prevent covered
22	persons and covered horses from using or
23	administering medications, substances, and
24	methods in violation of the rules estab-
25	lished in accordance with this Act;

1	(iv) exercise the powers specified in
2	section $6(c)(4)$ in accordance with that sec-
3	tion; and
4	(v) implement and undertake any
5	other responsibilities specified in the agree-
6	ment.
7	(F) TERM AND EXTENSION.—
8	(i) TERM OF INITIAL AGREEMENT
9	The initial agreement entered into by the
10	Authority under this paragraph shall be in
11	effect for the 5-year period beginning on
12	the program effective date.
13	(ii) EXTENSION.—At the end of the 5-
14	year period described in clause (i), the Au-
15	thority may—
16	(I) extend the term of the initial
17	agreement under this paragraph for
18	such additional term as is provided by
19	the rules of the Authority and con-
20	sistent with this Act; or
21	(II) enter into an agreement
22	meeting the requirements of this para-
23	graph with an entity described by sub-
24	paragraph (B) for such term as is

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provided by such rules and consistent
with this Act.
(2) AGREEMENTS FOR ENFORCEMENT BY
STATE RACING COMMISSIONS.—
(A) STATE RACING COMMISSIONS.—
(i) RACETRACK SAFETY PROGRAM.—
The Authority may enter into agreements
with State racing commissions for services
consistent with the enforcement of the
racetrack safety program.
(ii) ANTI-DOPING AND MEDICATION
CONTROL PROGRAM.—The anti-doping and
medication control enforcement agency
may enter into agreements with State rac-
ing commissions for services consistent
with the enforcement of the anti-doping
and medication control program.
(B) ELEMENTS OF AGREEMENTS.—Any
agreement under this paragraph shall include a
description of the scope of work, performance
metrics, reporting obligations, budgets, and any
other matter the Authority considers appro-
priate.
(3) Enforcement of standards.—The Au-
thority may coordinate with State racing commis-

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1	sions and other State regulatory agencies to monitor
2	and enforce racetrack compliance with the standards
3	developed under paragraphs (1) and (2) of section
4	7(e).
5	(f) PROCEDURES WITH RESPECT TO RULES OF AU-
6	THORITY.—
7	(1) ANTI-DOPING AND MEDICATION CON-
8	TROL.—
9	(A) IN GENERAL.—Recommendations for
10	rules regarding anti-doping and medication con-
11	trol shall be developed in accordance with sec-
12	tion 6.
13	(B) CONSULTATION.—The anti-doping and
14	medication control enforcement agency shall
15	consult with the anti-doping and medication
16	control standing committee and the Board of
17	the Authority on all anti-doping and medication
18	control rules of the Authority.
19	(2) RACETRACK SAFETY.—Recommendations
20	for rules regarding racetrack safety shall be devel-
21	oped by the racetrack safety standing committee of
22	the Authority.
23	(g) Subpoena and Investigatory Authority.—
24	The Authority shall have subpoena and investigatory au-

thority with respect to civil violations committed under its
 jurisdiction.

3 (h) CIVIL PENALTIES.—The Authority shall develop
4 a list of civil penalties with respect to the enforcement of
5 rules for covered persons and covered horseraces under its
6 jurisdiction.

7 (i) CIVIL ACTIONS.—

8 (1) IN GENERAL.—In addition to civil sanctions 9 imposed under section 8, the Authority may com-10 mence a civil action against a covered person or 11 racetrack that has engaged, is engaged, or is about 12 to engage, in acts or practices constituting a viola-13 tion of this Act or any rule established under this 14 Act in the proper district court of the United States, 15 the United States District Court for the District of 16 Columbia, or the United States courts of any terri-17 tory or other place subject to the jurisdiction of the 18 United States, to enjoin such acts or practices, to 19 enforce any civil sanctions imposed under that sec-20 tion, and for all other relief to which the Authority 21 may be entitled.

(2) INJUNCTIONS AND RESTRAINING ORDERS.—
With respect to a civil action commenced under
paragraph (1), upon a proper showing, a permanent

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1	or temporary injunction or restraining order shall be
2	granted without bond.
3	(j) Limitations on Authority.—
4	(1) PROSPECTIVE APPLICATION.—The jurisdic-
5	tion and authority of the Authority and the Commis-
6	sion with respect to the horseracing anti-doping and
7	medication control program and the racetrack safety
8	program shall be prospective only.
9	(2) Previous matters.—
10	(A) IN GENERAL.—The Authority and the
11	Commission may not investigate, prosecute, ad-
12	judicate, or penalize conduct in violation of the
13	horseracing anti-doping and medication control
14	program and the racetrack safety program that
15	occurs before the program effective date.
16	(B) STATE RACING COMMISSION.—With re-
17	spect to conduct described in subparagraph (A),
18	the applicable State racing commission shall re-
19	tain authority until the final resolution of the
20	matter.
21	(3) OTHER LAWS UNAFFECTED.—This Act
22	shall not be construed to modify, impair, or restrict
23	the operation of the general laws or regulations, as
24	may be amended from time to time, of the United
25	States, the States and their political subdivisions re-

lating to criminal conduct, cruelty to animals, mat ters unrelated to anti-doping, medication control and
 racetrack and racing safety of covered horses and
 covered races, and the use of medication in human
 participants in covered races.

6 (k) ELECTION FOR OTHER BREED COVERAGE7 UNDER ACT.—

8 (1) IN GENERAL.—A State racing commission 9 or a breed governing organization for a breed of 10 horses other than Thoroughbred horses may elect to 11 have such breed be covered by this Act by the filing 12 of a designated election form and subsequent ap-13 proval by the Authority. A State racing commission 14 may elect to have a breed covered by this Act for the 15 applicable State only.

16 (2)ELECTION CONDITIONAL ON FUNDING 17 MECHANISM.—A commission or organization may 18 not make an election under paragraph (1) unless the 19 commission or organization has in place a mecha-20 nism to provide sufficient funds to cover the costs of 21 the administration of this Act with respect to the 22 horses that will be covered by this Act as a result 23 of the election.

24 (3) APPORTIONMENT.—The Authority shall apportion costs described in paragraph (2) in connec-

tion with an election under paragraph (1) fairly
among all impacted segments of the horseracing industry, subject to approval by the Commission in accordance with section 4. Such apportionment may
not provide for the allocation of costs or funds
among breeds of horses.

7 SEC. 6. HORSERACING ANTI-DOPING AND MEDICATION 8 CONTROL PROGRAM.

9 (a) PROGRAM REQUIRED.—

10 (1) IN GENERAL.—Not later than the program 11 effective date, and after notice and an opportunity 12 for public comment in accordance with section 4, the 13 Authority shall establish a horseracing anti-doping 14 and medication control program applicable to all 15 covered horses, covered persons, and covered 16 horseraces in accordance with the registration of 17 covered persons under section 5(d).

18 (2) Consideration of other breeds.—In 19 developing the horseracing anti-doping and medica-20 tion control program with respect to a breed of horse 21 that is made subject to this Act by election of a 22 State racing commission or the breed governing or-23 ganization for such horse under section 5(k), the 24 Authority shall consider the unique characteristics of 25 such breed.

1 (b) CONSIDERATIONS IN DEVELOPMENT OF PRO-2 GRAM.—In developing the horseracing anti-doping and 3 medication control program, the Authority shall take into 4 consideration the following:

5 (1) Covered horses should compete only when
6 they are free from the influence of medications,
7 other foreign substances, and methods that affect
8 their performance.

9 (2) Covered horses that are injured or unsound 10 should not train or participate in covered races, and 11 the use of medications, other foreign substances, and 12 treatment methods that mask or deaden pain in 13 order to allow injured or unsound horses to train or 14 race should be prohibited.

(3) Rules, standards, procedures, and protocols
regulating medication and treatment methods for
covered horses and covered races should be uniform
and uniformly administered nationally.

(4) To the extent consistent with this Act, consideration should be given to international antidoping and medication control standards of the
International Federation of Horseracing Authorities
and the Principles of Veterinary Medical Ethics of
the American Veterinary Medical Association.

1 (5) The administration of medications and 2 treatment methods to covered horses should be 3 based upon an examination and diagnosis that iden-4 tifies an issue requiring treatment for which the 5 medication or method represents an appropriate 6 component of treatment.

7 (6) The amount of therapeutic medication that
8 a covered horse receives should be the minimum nec9 essary to address the diagnosed health concerns
10 identified during the examination and diagnostic
11 process.

(7) The welfare of covered horses, the integrity
of the sport, and the confidence of the betting public
require full disclosure to regulatory authorities regarding the administration of medications and treatments to covered horses.

17 (c) ACTIVITIES.—The following activities shall be car18 ried out under the horseracing anti-doping and medication
19 control program:

(1) STANDARDS FOR ANTI-DOPING AND MEDICATION CONTROL.—Not later than 120 days before
the program effective date, the Authority shall issue,
by rule—

24 (A) uniform standards for—

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(i) the administration of medication to
covered horses by covered persons; and
(ii) laboratory testing accreditation
and protocols; and
(B) a list of permitted and prohibited
medications, substances, and methods, including
allowable limits of permitted medications, sub-
stances, and methods.
(2) Review process for administration of
MEDICATION.—The development of a review process
for the administration of any medication to a cov-
ered horse during the 48-hour period preceding the
next racing start of the covered horse.
(3) AGREEMENT REQUIREMENTS.—The devel-
opment of requirements with respect to agreements
under section 5(e).
(4) ANTI-DOPING AND MEDICATION CONTROL
ENFORCEMENT AGENCY.—
(A) CONTROL RULES, PROTOCOLS, ETC.—
Except as provided in paragraph (5), the anti-
doping and medication control program enforce-
ment agency under section 5(e) shall, in con-
sultation with the anti-doping and medication
control standing committee of the Authority
and consistent with international best practices,

develop and recommend anti-doping and medication control rules, protocols, policies, and guidelines for approval by the Authority.

(B) RESULTS MANAGEMENT.—The anti-4 doping and medication control enforcement 5 6 agency shall conduct and oversee anti-doping 7 and medication control results management, in-8 cluding independent investigations, charging 9 and adjudication of potential medication control 10 rule violations, and the enforcement of any civil 11 sanctions for such violations. Any final decision 12 or civil sanction of the anti-doping and medica-13 tion control enforcement agency under this sub-14 paragraph shall be the final decision or civil 15 sanction of the Authority, subject to review in 16 accordance with section 9.

17 (C) TESTING.—The anti-doping enforce18 ment agency shall perform and manage test dis19 tribution planning (including intelligence-based
20 testing), the sample collection process, and in21 competition and out-of-competition testing (in22 cluding no-advance-notice testing).

23 (D) TESTING LABORATORIES.—The anti24 doping and medication control enforcement
25 agency shall accredit testing laboratories based

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upon the standards established under this Act, and shall monitor, test, and audit accredited laboratories to ensure continuing compliance with accreditation standards.

5 (5) ANTI-DOPING AND MEDICATION CONTROL 6 STANDING COMMITTEE.—The anti-doping and medi-7 cation control standing committee shall, in consulta-8 tion with the anti-doping and medication control en-9 forcement agency, develop lists of permitted and pro-10 hibited medications, methods, and substances for 11 recommendation to, and approval by, the Authority. 12 Any such list may prohibit the administration of any 13 substance or method to a horse at any time after 14 such horse becomes a covered horse if the Authority 15 determines such substance or method has a long-16 term degrading effect on the soundness of a horse. 17 (d) PROHIBITION.—Except as provided in sub-18 sections (e) and (f), the horseracing anti-doping and medication control program shall prohibit the administration 19 20 of any prohibited or otherwise permitted substance to a 21 covered horse within 48 hours of its next racing start, ef-22 fective as of the program effective date.

23 (e) Advisory Committee Study and Report.—

24 (1) IN GENERAL.—Not later than the program
25 effective date, the Authority shall convene an advi-

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1	sory committee comprised of horseracing anti-doping
2	and medication control industry experts, including a
3	member designated by the anti-doping and medica-
4	tion control enforcement agency, to conduct a study
5	on the use of furosemide on horses during the 48-
6	hour period before the start of a race, including the
7	effect of furosemide on equine health and the integ-
8	rity of competition and any other matter the Author-
9	ity considers appropriate.
10	(2) Report.—Not later than three years after
11	the program effective date, the Authority shall direct
12	the advisory committee convened under paragraph
13	(1) to submit to the Authority a written report on
14	the study conducted under that paragraph that in-
15	cludes recommended changes, if any, to the prohibi-
16	tion in subsection (d).
17	(3) Modification of prohibition.—
18	(A) IN GENERAL.—After receipt of the re-
19	port required by paragraph (2), the Authority
20	may, by unanimous vote of the Board of the
21	Authority, modify the prohibition in subsection
22	(d) and, notwithstanding subsection (f), any
23	such modification shall apply to all States be-
24	ginning on the date that is three years after the
25	program effective date.

1	(B) CONDITION.—In order for a unani-
2	mous vote described in subparagraph (A) to ef-
3	fect a modification of the prohibition in sub-
4	section (d), the vote must include unanimous
5	adoption of each of the following findings:
6	(i) That the modification is war-
7	ranted.
8	(ii) That the modification is in the
9	best interests of horse racing.
10	(iii) That furosemide has no perform-
11	ance enhancing effect on individual horses.
12	(iv) That public confidence in the in-
13	tegrity and safety of racing would not be
14	adversely affected by the modification.
15	(f) EXEMPTION.—
16	(1) IN GENERAL.—Except as provided in para-
17	graph (2), only during the three-year period begin-
18	ning on the program effective date, a State racing
19	commission may submit to the Authority, at such
20	time and in such manner as the Authority may re-
21	quire, a request for an exemption from the prohibi-
22	tion in subsection (d) with respect to the use of
23	furosemide on covered horses during such period.
24	(2) EXCEPTIONS.—An exemption under para-
25	graph (1) may not be requested for—

1	(A) two-year-old covered horses; or
2	(B) covered horses competing in stakes
3	races.
4	(3) CONTENTS OF REQUEST.—A request under
5	paragraph (1) shall specify the applicable State rac-
6	ing commission's requested limitations on the use of
7	furosemide that would apply to the State under the
8	horseracing anti-doping and medication control pro-
9	gram during such period. Such limitations shall be
10	no less restrictive on the use and administration of
11	furosemide than the restrictions set forth in State's
12	laws and regulations in effect as of September 1,
13	2020.

(4) GRANT OF EXEMPTION.—Subject to subsection (e)(3), the Authority shall grant an exemption requested under paragraph (1) for the remainder of such period and shall allow the use of
furosemide on covered horses in the applicable State,
in accordance with the requested limitations.

20 (g) BASELINE ANTI-DOPING AND MEDICATION CON-21 TROL RULES.—

(1) IN GENERAL.—Subject to paragraph (3),
the baseline anti-doping and medication control rules
described in paragraph (2) shall—

1	(A) constitute the initial rules of the horse-
2	racing anti-doping and medication control pro-
3	gram; and
4	(B) except as exempted pursuant to sub-
5	sections (e) and (f), remain in effect at all
6	times after the program effective date.
7	(2) BASELINE ANTI-DOPING MEDICATION CON-
8	TROL RULES DESCRIBED.—
9	(A) IN GENERAL.—The baseline anti-
10	doping and medication control rules described
11	in this paragraph are the following:
12	(i) The lists of permitted and prohib-
13	ited substances (including drugs, medica-
14	tions, and naturally occurring substances
15	and synthetically occurring substances) in
16	effect for the International Federation of
17	Horseracing Authorities, including the
18	International Federation of Horseracing
19	Authorities International Screening Limits
20	for urine, dated May 2019, and the Inter-
21	national Federation of Horseracing Au-
22	thorities International Screening Limits for
23	plasma, dated May 2019.

1	(ii) The World Anti-Doping Agency
2	International Standard for Laboratories
3	(version 10.0), dated November 12, 2019.
4	(iii) The Association of Racing Com-
5	missioners International out-of-competition
6	testing standards, Model Rules of Racing
7	(version 9.2).
8	(iv) The Association of Racing Com-
9	missioners International penalty and mul-
10	tiple medication violation rules, Model
11	Rules of Racing (version 6.2).
12	(B) CONFLICT OF RULES.—In the case of
13	a conflict among the rules described in subpara-
14	graph (A), the most stringent rule shall apply.
15	(3) Modifications to baseline rules.—
16	(A) DEVELOPMENT BY ANTI-DOPING AND
17	MEDICATION CONTROL STANDING COM-
18	MITTEE.—The anti-doping and medication con-
19	trol standing committee, in consultation with
20	the anti-doping and medication control enforce-
21	ment agency, may develop and submit to the
22	Authority for approval by the Authority pro-
23	posed modifications to the baseline anti-doping
24	and medication control rules.

1 (B) AUTHORITY APPROVAL.—If the Au-2 thority approves a proposed modification under 3 this paragraph, the proposed modification shall 4 be submitted to and considered by the Commis-5 sion in accordance with section 4. 6 (C) ANTI-DOPING AND MEDICATION CON-7 TROL ENFORCEMENT AGENCY VETO AUTHOR-8 ITY.—The Authority shall not approve any pro-9 posed modification that renders an anti-doping 10 and medication control rule less stringent than 11 the baseline anti-doping and medication control 12 rules described in paragraph (2) (including by 13 increasing permitted medication thresholds. 14 adding permitted medications, removing prohib-15 ited medications, or weakening enforcement 16 mechanisms) without the approval of the anti-17 doping and medication control enforcement 18 agency.

19 SEC. 7. RACETRACK SAFETY PROGRAM.

20 (a) Establishment and Considerations.—

(1) IN GENERAL.—Not later than the program
effective date, and after notice and an opportunity
for public comment in accordance with section 4, the
Authority shall establish a racetrack safety program
applicable to all covered horses, covered persons, and

covered horseraces in accordance with the registra tion of covered persons under section 5(d).

3 (2)CONSIDERATIONS IN DEVELOPMENT OF 4 SAFETY PROGRAM.—In the development of the 5 horseracing safety program for covered horses, cov-6 ered persons, and covered horseraces, the Authority 7 and the Commission shall take into consideration ex-8 isting safety standards including the National Thor-9 oughbred Racing Association Safety and Integrity 10 Alliance Code of Standards, the International Fed-11 eration of Horseracing Authority's International 12 Agreement on Breeding, Racing, and Wagering, and 13 the British Horseracing Authority's Equine Health 14 and Welfare program.

(b) ELEMENTS OF HORSERACING SAFETY PRO16 GRAM.—The horseracing safety program shall include the
17 following:

18 (1) A set of training and racing safety stand19 ards and protocols taking into account regional dif20 ferences and the character of differing racing facili21 ties.

(2) A uniform set of training and racing safety
standards and protocols consistent with the humane
treatment of covered horses, which may include lists

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1	of permitted and prohibited practices or methods
2	(such as crop use).
3	(3) A racing surface quality maintenance sys-
4	tem that—
5	(A) takes into account regional differences
6	and the character of differing racing facilities;
7	and
8	(B) may include requirements for track
9	surface design and consistency and established
10	standard operating procedures related to track
11	surface, monitoring, and maintenance (such as
12	standardized seasonal assessment, daily track-
13	ing, and measurement).
14	(4) A uniform set of track safety standards and
15	protocols, that may include rules governing oversight
16	and movement of covered horses and human and
17	equine injury reporting and prevention.
18	(5) Programs for injury and fatality data anal-
19	ysis, that may include pre- and post-training and
20	race inspections, use of a veterinarian's list, and
21	concussion protocols.
22	(6) The undertaking of investigations at race-
23	track and non-racetrack facilities related to safety
24	violations.

1	(7) Procedures for investigating, charging, and
2	adjudicating violations and for the enforcement of
3	civil sanctions for violations.
4	(8) A schedule of civil sanctions for violations.
5	(9) Disciplinary hearings, which may include
6	binding arbitration, civil sanctions, and research.
7	(10) Management of violation results.
8	(11) Programs relating to safety and perform-
9	ance research and education.
10	(12) An evaluation and accreditation program
11	that ensures that racetracks in the United States
12	meet the standards described in the elements of the
13	Horseracing Safety Program.
14	(c) ACTIVITIES.—The following activities shall be car-
15	ried out under the racetrack safety program:
16	(1) STANDARDS FOR RACETRACK SAFETY.—
17	The development, by the racetrack safety standing
18	committee of the Authority in section $3(c)(2)$ of uni-
19	form standards for racetrack and horseracing safety.
20	(2) STANDARDS FOR SAFETY AND PERFORM-
21	ANCE ACCREDITATION.—
22	(A) IN GENERAL.—Not later than 120
23	days before the program effective date, the Au-
24	thority, in consultation with the racetrack safe-

1	ty standing committee, shall issue, by rule in
2	accordance with section 4—
3	(i) safety and performance standards
4	of accreditation for racetracks; and
5	(ii) the process by which a racetrack
6	may achieve and maintain accreditation by
7	the Authority.
8	(B) Modifications.—
9	(i) IN GENERAL.—The Authority may
10	modify rules establishing the standards
11	issued under subparagraph (A), as the Au-
12	thority considers appropriate.
13	(ii) NOTICE AND COMMENT.—The
14	Commission shall publish in the Federal
15	Register any proposed rule of the Author-
16	ity, and provide an opportunity for public
17	comment with respect to, any modification
18	under clause (i) in accordance with section
19	4.
20	(C) EXTENSION OF PROVISIONAL OR IN-
21	TERIM ACCREDITATION.—The Authority may,
22	by rule in accordance with section 4, extend
23	provisional or interim accreditation to a race-
24	track accredited by the National Thoroughbred
25	Racing Association Safety and Integrity Alli-

ance on a date before the program effective
 date.

3 (3) NATIONWIDE SAFETY AND PERFORMANCE
4 DATABASE.—

(A) IN GENERAL.—Not later than one year 5 6 after the program effective date, and after no-7 tice and an opportunity for public comment in 8 accordance with section 4, the Authority, in 9 consultation with the Commission, shall develop 10 and maintain a nationwide database of race-11 horse safety, performance, health, and injury 12 information for the purpose of conducting an 13 epidemiological study.

14 (B) COLLECTION OF INFORMATION.—In 15 accordance with the registration of covered per-16 sons under section 5(d), the Authority may re-17 quire covered persons to collect and submit to 18 the database described in subparagraph (A) 19 such information as the Authority may require 20 to further the goal of increased racehorse wel-21 fare.

22 SEC. 8. RULE VIOLATIONS AND CIVIL SANCTIONS.

23 (a) DESCRIPTION OF RULE VIOLATIONS.—

24 (1) IN GENERAL.—The Authority shall issue, by
25 rule in accordance with section 4, a description of

1	safety, performance, and anti-doping and medication
2	control rule violations applicable to covered horses
3	and covered persons.
4	(2) Elements.—The description of rule viola-
5	tions established under paragraph (1) may include
6	the following:
7	(A) With respect to a covered horse, strict
8	liability for covered trainers for—
9	(i) the presence of a prohibited sub-
10	stance or method in a sample or the use of
11	a prohibited substance or method;
12	(ii) the presence of a permitted sub-
13	stance in a sample in excess of the amount
14	allowed by the horseracing anti-doping and
15	medication control program; and
16	(iii) the use of a permitted method in
17	violation of the applicable limitations es-
18	tablished under the horseracing anti-
19	doping and medication control program.
20	(B) Attempted use of a prohibited sub-
21	stance or method on a covered horse.
22	(C) Possession of any prohibited substance
23	or method.
24	(D) Attempted possession of any prohib-
25	ited substance or method.

(E) Administration or attempted adminis-
tration of any prohibited substance or method
on a covered horse.
(F) Refusal or failure, without compelling
justification, to submit a covered horse for sam-
ple collection.
(G) Failure to cooperate with the Author-
ity or an agent of the Authority during any in-
vestigation.
(H) Failure to respond truthfully, to the
best of a covered person's knowledge, to a ques-
tion of the Authority or an agent of the Author-
ity with respect to any matter under the juris-
diction of the Authority.
(I) Tampering or attempted tampering
with the application of the safety, performance,
or anti-doping and medication control rules or
process adopted by the Authority, including—
(i) the intentional interference, or an
attempt to interfere, with an official or
agent of the Authority;
(ii) the procurement or the provision
of fraudulent information to the Authority
or agent; and

1	(iii) the intimidation of, or an attempt
2	to intimidate, a potential witness.
3	(J) Trafficking or attempted trafficking in
4	any prohibited substance or method.
5	(K) Assisting, encouraging, aiding, abet-
6	ting, conspiring, covering up, or any other type
7	of intentional complicity involving a safety, per-
8	formance, or anti-doping and medication control
9	rule violation or the violation of a period of sus-
10	pension or eligibility.
11	(L) Threatening or seeking to intimidate a
12	person with the intent of discouraging the per-
13	son from the good faith reporting to the Au-
14	thority, an agent of the Authority or the Com-
15	mission, or the anti-doping and medication con-
16	trol enforcement agency under section 5(e), of
17	information that relates to—
18	(i) an alleged safety, performance, or
19	anti-doping and medication control rule
20	violation; or
21	(ii) alleged noncompliance with a safe-
22	ty, performance, or anti-doping and medi-
23	cation control rule.
24	(b) Testing Laboratories.—

1	(1) Accreditation and standards.—Not
2	later than 120 days before the program effective
3	date, the Authority shall, in consultation with the
4	anti-doping and medication control enforcement
5	agency, establish, by rule in accordance with section
6	4—
7	(A) standards of accreditation for labora-
8	tories involved in testing samples from covered
9	horses;
10	(B) the process for achieving and main-
11	taining accreditation; and
12	(C) the standards and protocols for testing
10	an de generales
13	such samples.
13 14	(2) ADMINISTRATION.—The accreditation of
	*
14	(2) ADMINISTRATION.—The accreditation of
14 15	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited
14 15 16	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority
14 15 16 17	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority rules shall be administered by the anti-doping and
14 15 16 17 18	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority rules shall be administered by the anti-doping and medication control enforcement agency. The anti-
14 15 16 17 18 19	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority rules shall be administered by the anti-doping and medication control enforcement agency. The anti- doping and medication control enforcement agency
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14 15 16 17 18 19 20 21	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority rules shall be administered by the anti-doping and medication control enforcement agency. The anti- doping and medication control enforcement agency shall have the authority to require specific test sam- ples to be directed to and tested by laboratories hav-
 14 15 16 17 18 19 20 21 22 	(2) ADMINISTRATION.—The accreditation of laboratories and the conduct of audits of accredited laboratories to ensure compliance with Authority rules shall be administered by the anti-doping and medication control enforcement agency. The anti- doping and medication control enforcement agency shall have the authority to require specific test sam- ples to be directed to and tested by laboratories hav- ing special expertise in the required tests.

1	toring accorditation to a laboratory accordited by the
1	terim accreditation to a laboratory accredited by the
2	Racing Medication and Testing Consortium, Inc., on
3	a date before the program effective date.
4	(4) Selection of Laboratories.—
5	(A) IN GENERAL.—Except as provided in
6	paragraph (2), a State racing commission may
7	select a laboratory accredited in accordance
8	with the standards established under paragraph
9	(1) to test samples taken in the applicable
10	State.
11	(B) SELECTION BY THE AUTHORITY.—If a
12	State racing commission does not select an ac-
13	credited laboratory under subparagraph (A),
14	the Authority shall select such a laboratory to
15	test samples taken in the State concerned.
16	(c) Results Management and Disciplinary
17	PROCESS.—
18	(1) IN GENERAL.—Not later than 120 days be-
19	fore the program effective date, the Authority shall
20	establish in accordance with section 4—
21	(A) rules for safety, performance, and anti-
22	doping and medication control results manage-
23	ment; and

1	(B) the disciplinary process for safety, per-
2	formance, and anti-doping and medication con-
3	trol rule violations.
4	(2) ELEMENTS.—The rules and process estab-
5	lished under paragraph (1) shall include the fol-
6	lowing:
7	(A) Provisions for notification of safety,
8	performance, and anti-doping and medication
9	control rule violations.
10	(B) Hearing procedures.
11	(C) Standards for burden of proof.
12	(D) Presumptions.
13	(E) Evidentiary rules.
14	(F) Appeals.
15	(G) Guidelines for confidentiality and pub-
16	lic reporting of decisions.
17	(3) DUE PROCESS.—The rules established
18	under paragraph (1) shall provide for adequate due
19	process, including impartial hearing officers or tribu-
20	nals commensurate with the seriousness of the al-
21	leged safety, performance, or anti-doping and medi-
22	cation control rule violation and the possible civil
23	sanctions for such violation.
24	(d) Civil Sanctions.—

1	(1) IN GENERAL.—The Authority shall estab-
2	lish uniform rules, in accordance with section 4, im-
3	posing civil sanctions against covered persons or cov-
4	ered horses for safety, performance, and anti-doping
5	and medication control rule violations.
6	(2) REQUIREMENTS.—The rules established
7	under paragraph (1) shall—
8	(A) take into account the unique aspects of
9	horseracing;
10	(B) be designed to ensure fair and trans-
11	parent horseraces; and
12	(C) deter safety, performance, and anti-
13	doping and medication control rule violations.
14	(3) Severity.—The civil sanctions under para-
15	graph (1) may include—
16	(A) lifetime bans from horseracing,
17	disgorgement of purses, monetary fines and
18	penalties, and changes to the order of finish in
19	covered races; and
20	(B) with respect to anti-doping and medi-
21	cation control rule violators, an opportunity to
22	reduce the applicable civil sanctions that is
23	comparable to the opportunity provided by the
24	Protocol for Olympic Movement Testing of the
25	United States Anti-Doping Agency.

(e) MODIFICATIONS.—The Authority may propose a
 modification to any rule established under this section as
 the Authority considers appropriate, and the proposed
 modification shall be submitted to and considered by the
 Commission in accordance with section 4.

6 SEC. 9. REVIEW OF FINAL DECISIONS OF THE AUTHORITY.

7 (a) NOTICE OF CIVIL SANCTIONS.—If the Authority
8 imposes a final civil sanction for a violation committed by
9 a covered person pursuant to the rules or standards of
10 the Authority, the Authority shall promptly submit to the
11 Commission notice of the civil sanction in such form as
12 the Commission may require.

13 (b) REVIEW BY ADMINISTRATIVE LAW JUDGE.—

14 (1) IN GENERAL.—With respect to a final civil
15 sanction imposed by the Authority, on application by
16 the Commission or a person aggrieved by the civil
17 sanction filed not later than 30 days after the date
18 on which notice under subsection (a) is submitted,
19 the civil sanction shall be subject to de novo review
20 by an administrative law judge.

21 (2) NATURE OF REVIEW.—

22 (A) IN GENERAL.—In matters reviewed
23 under this subsection, the administrative law
24 judge shall determine whether—

1	(i) a person has engaged in such acts
2	or practices, or has omitted such acts or
3	practices, as the Authority has found the
4	person to have engaged in or omitted;
5	(ii) such acts, practices, or omissions
6	are in violation of this Act or the anti-
7	doping and medication control or racetrack
8	safety rules approved by the Commission;
9	or
10	(iii) the final civil sanction of the Au-
11	thority was arbitrary, capricious, an abuse
12	of discretion, or otherwise not in accord-
13	ance with law.
14	(B) CONDUCT OF HEARING.—An adminis-
15	trative law judge shall conduct a hearing under
16	this subsection in such a manner as the Com-
17	mission may specify by rule, which shall con-
18	form to section 556 of title 5, United States
19	Code.
20	(3) Decision by administrative law
21	JUDGE.—
22	(A) IN GENERAL.—With respect to a mat-
23	ter reviewed under this subsection, an adminis-
24	trative law judge—

- 1 (i) shall render a decision not later than 60 days after the conclusion of the 2 3 hearing; 4 (ii) may affirm, reverse, modify, set 5 aside, or remand for further proceedings, 6 in whole or in part, the final civil sanction 7 of the Authority; and 8 (iii) may make any finding or conclu-9 sion that, in the judgment of the adminis-10 trative law judge, is proper and based on 11 the record. 12 (B) FINAL DECISION.—A decision under 13 this paragraph shall constitute the decision of 14 the Commission without further proceedings 15 unless a notice or an application for review is 16 timely filed under subsection (c). 17 (c) REVIEW BY COMMISSION.— 18 (1) NOTICE OF REVIEW BY COMMISSION.—The 19 Commission may, on its own motion, review any de-20 cision of an administrative law judge issued under subsection (b)(3) by providing written notice to the 21 22 Authority and any interested party not later than 30 23 days after the date on which the administrative law 24 judge issues the decision.
- 25 (2) Application for review.—

(A) IN GENERAL.—The Authority or a per-
son aggrieved by a decision issued under sub-
section $(b)(3)$ may petition the Commission for
review of such decision by filing an application
for review not later than 30 days after the date
on which the administrative law judge issues
the decision.
(B) EFFECT OF DENIAL OF APPLICATION
FOR REVIEW.—If an application for review
under subparagraph (A) is denied, the decision
of the administrative law judge shall constitute
the decision of the Commission without further
proceedings.
(C) DISCRETION OF COMMISSION.—
(i) IN GENERAL.—A decision with re-
spect to whether to grant an application
for review under subparagraph (A) is sub-
ject to the discretion of the Commission.
(ii) Matters to be considered.—
In determining whether to grant such an
application for review, the Commission
shall consider whether the application
makes a reasonable showing that—

	01
1	(I) a prejudicial error was com-
2	mitted in the conduct of the pro-
3	ceeding; or
4	(II) the decision involved—
5	(aa) an erroneous applica-
6	tion of the anti-doping and medi-
7	cation control or racetrack safety
8	rules approved by the Commis-
9	sion; or
10	(bb) an exercise of discretion
11	or a decision of law or policy that
12	warrants review by the Commis-
13	sion.
14	(3) NATURE OF REVIEW.—
15	(A) IN GENERAL.—In matters reviewed
16	under this subsection, the Commission may—
17	(i) affirm, reverse, modify, set aside,
18	or remand for further proceedings, in
19	whole or in part, the decision of the admin-
20	istrative law judge; and
21	(ii) make any finding or conclusion
22	that, in the judgement of the Commission,
23	is proper and based on the record.
24	(B) DE NOVO REVIEW.—The Commission
25	shall review de novo the factual findings and

	<u> </u>
1	conclusions of law made by the administrative
2	law judge.
3	(C) Consideration of additional evi-
4	DENCE.—
5	(i) MOTION BY COMMISSION.—The
6	Commission may, on its own motion, allow
7	the consideration of additional evidence.
8	(ii) MOTION BY A PARTY.—
9	(I) IN GENERAL.—A party may
10	file a motion to consider additional
11	evidence at any time before the
12	issuance of a decision by the Commis-
13	sion, which shall show, with particu-
14	larity, that—
15	(aa) such additional evidence
16	is material; and
17	(bb) there were reasonable
18	grounds for failure to submit the
19	evidence previously.
20	(II) PROCEDURE.—The Commis-
21	sion may—
22	(aa) accept or hear addi-
23	tional evidence; or
24	(bb) remand the proceeding
25	to the administrative law judge

1	for	the	consideration	of	addi-
2	tional evidence.				

3 (d) STAY OF PROCEEDINGS.—Review by an adminis4 trative law judge or the Commission under this section
5 shall not operate as a stay of a final civil sanction of the
6 Authority unless the administrative law judge or Commis7 sion orders such a stay.

8 SEC. 10. UNFAIR OR DECEPTIVE ACTS OR PRACTICES.

9 The sale of a covered horse, or of any other horse 10 in anticipation of its future participation in a covered race, shall be considered an unfair or deceptive act or practice 11 in or affecting commerce under section 5(a) of the Federal 12 13 Trade Commission Act (15 U.S.C. 45(a)) if the seller— 14 (1) knows or has reason to know the horse has 15 been administered— 16 (A) a bisphosphonate prior to the horse's 17 fourth birthday; or 18 (B) any other substance or method the Au-19 thority determines has a long-term degrading 20 effect on the soundness of the covered horse; 21 and 22 (2) fails to disclose to the buyer the administra-23 tion of the bisphosphonate or other substance or 24 method described in paragraph (1)(B).

1 SEC. 11. STATE DELEGATION; COOPERATION.

2 (a) STATE DELEGATION.—

3 (1) IN GENERAL.—The Authority may enter 4 into an agreement with a State racing commission to 5 implement, within the jurisdiction of the State rac-6 ing commission, a component of the racetrack safety 7 program or, with the concurrence of the anti-doping 8 and medication control enforcement agency under section 5(e), a component of the horseracing anti-9 10 doping and medication control program, if the Au-11 thority determines that the State racing commission 12 has the ability to implement such component in ac-13 cordance with the rules, standards, and require-14 ments established by the Authority.

(2) IMPLEMENTATION BY STATE RACING COMMISSION.—A State racing commission or other appropriate regulatory body of a State may not implement such a component in a manner less restrictive
than the rule, standard, or requirement established
by the Authority.

(b) COOPERATION.—To avoid duplication of functions, facilities, and personnel, and to attain closer coordination and greater effectiveness and economy in administration of Federal and State law, where conduct by any
person subject to the horseracing medication control program or the racetrack safety program may involve both

a medication control or racetrack safety rule violation and
 violation of Federal or State law, the Authority and Fed eral or State law enforcement authorities shall cooperate
 and share information.