AM	IENDMENT NO Calendar No
Pu	rpose: In the nature of a substitute.
IN	THE SENATE OF THE UNITED STATES—118th Cong., 2d Sess.
	S. 4667
То	amend title 31, United States Code, to establish the Life Sciences Research Security Board, and for other purposes.
R	eferred to the Committee on and ordered to be printed
	Ordered to lie on the table and to be printed
A	MENDMENT IN THE NATURE OF A SUBSTITUTE intended to be proposed by Mr. Paul
Viz	:
1	Strike all after the enacting clause and insert the fol-
2	lowing:
3	SECTION 1. SHORT TITLE.
4	This Act may be cited as the "Risky Research Review
5	Act".
6	SEC. 2. LIFE SCIENCES RESEARCH SECURITY BOARD.
7	(a) In General.—Subtitle V of title 31, United
8	States Code, is amended by adding at the end the fol-
9	lowing:

1	"CHAPTER 79—LIFE SCIENCES RESEARCH
2	SECURITY BOARD
3	"§ 7901. Definitions
4	"In this chapter:
5	"(1) Agency.—The term 'agency' has the
6	meaning given the term in section 552(f) of title 5
7	"(2) Appropriate congressional commit-
8	TEES.—The term 'appropriate congressional com-
9	mittees' means the Committee on Homeland Secu-
10	rity and Governmental Affairs of the Senate and the
11	Committee on Oversight and Accountability of the
12	House of Representatives.
13	"(3) Board.—The term 'Board' means the
14	Life Sciences Research Security Board established
15	under section 7902(a).
16	"(4) Dual use research of concern.—The
17	term 'dual use research of concern'—
18	"(A) means life sciences research that
19	based on current understanding, can be reason-
20	ably anticipated to provide knowledge, informa-
21	tion, products, or technologies that could—
22	"(i) be misapplied to do harm with no
23	modification or only a minor modification
24	and

1	"(ii) pose a significant threat with po-
2	tential consequences to public health and
3	safety, agricultural crops and other plants,
4	animals, materiel, or national security; and
5	"(B) includes—
6	"(i) life sciences research that could—
7	"(I) increase transmissibility of a
8	pathogen within or between host spe-
9	cies;
10	"(II) increase the virulence of a
11	pathogen or convey virulence to a non-
12	pathogen;
13	"(III) increase the toxicity of a
14	known toxin or produce a novel toxin;
15	"(IV) increase—
16	"(aa) the stability of a
17	pathogen or toxin in the environ-
18	ment; or
19	"(bb) the ability to dissemi-
20	nate a pathogen or toxin;
21	"(V) alter the host range or tro-
22	pism of a pathogen or toxin;
23	"(VI) decrease the ability for a
24	human or veterinary pathogen or

1	toxin to be detected using standard
2	diagnostic or analytical methods;
3	"(VII) increase resistance of a
4	pathogen or toxin to clinical or veteri-
5	nary prophylactic or therapeutic inter-
6	ventions;
7	"(VIII) alter a human or veteri-
8	nary pathogen or toxin to disrupt the
9	effectiveness of pre-existing immunity,
10	via immunization or natural infection,
11	against the pathogen or toxin;
12	"(IX) enhance the susceptibility
13	of a host population to a pathogen or
14	toxin;
15	"(X) enhance transmissibility of
16	a pathogen in humans;
17	"(XI) enhance the virulence of a
18	pathogen in humans;
19	"(XII) enhance the immune eva-
20	sion of a pathogen in humans, such as
21	by modifying the pathogen to disrupt
22	the effectiveness of pre-existing immu-
23	nity via immunization or natural in-
24	fection; or

1	"(XIII) generate, use, reconsti-
2	tute, or transfer an eradicated or ex-
3	tinct high-consequence pathogen; and
4	"(ii) any other category of life
5	sciences research that the Board, by ma-
6	jority vote of the members of the Board,
7	identifies and publishes in the Federal
8	Register.
9	"(5) Employee.—The term 'employee' means
10	an individual described in section 2105(a) of title 5.
11	"(6) Federal funding.—The term 'Federal
12	funding' means amounts awarded by an agency pur-
13	suant to an intramural or extramural grant, cooper-
14	ative agreement, interagency agreement, contract, or
15	other instrument.
16	"(7) Gain of function research.—The
17	term 'gain of function research' means a research
18	experiment that may enhance the transmissibility or
19	virulence of a high-consequence pathogen.
20	"(8) High-consequence pathogen.—The
21	term 'high-consequence pathogen'—
22	"(A) means a wild-type or synthetic patho-
23	gen that—

1	"(i)(I) is likely capable of wide and
2	uncontrollable spread in human popu-
3	lations; and
4	"(II) would likely cause moderate to
5	severe disease or mortality in humans; or
6	"(ii) is—
7	"(I) subject to subparagraph (B),
8	influenza A virus;
9	"(II) classified under subgenus
10	Sarbecovirus;
11	"(III) classified under subgenus
12	Merbecovirus;
13	"(IV) Variola orthopoxvirus;
14	"(V) Mpox orthopoxvirus;
15	"(VI) Nipah henipavirus;
16	"(VII) Hendra henipavirus;
17	"(VIII) Ebola orthoebolavirus;
18	"(IX) Marburg marburgvirus
19	"(X) Lassa mammarenavirus;
20	"(XI) Junin arenavirus;
21	"(XII) Crimean-Congo hemor-
22	rhagic fever orthonairovirus;
23	"(XIII) Hantaan
24	orthohantavirus;

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1	"(XIV) Sin Nombre
2	orthohantavirus;
3	"(XV) Yersinia pestis;
4	"(XVI) a select agent or toxin
5	work with which poses a significant
6	risk of deliberate misuse;
7	"(XVII) any other pathogen or
8	category of pathogen that a majority
9	of members of the Board—
10	"(aa) identifies as a high-
11	consequence pathogen; and
12	"(bb) publishes in the Fed-
13	eral Register; or
14	"(XVIII) any synthetic construct
15	of a pathogen or category of pathogen
16	described in this clause; and
17	"(B) does not include a seasonal influenza
18	virus, unless a seasonal influenza virus has been
19	manipulated to include genetic sequences from
20	a pathogen described in subparagraph (A).
21	"(9) High-risk life sciences research.—
22	The term 'high-risk life sciences research' means life
23	sciences research that is—
24	"(A) dual use research of concern involving
25	a high-consequence pathogen; or

"(A) means the study or use of a living ganism, a virus, or a product of a living or nism or virus; and "(B) includes each discipline, methodology and application of biology, including by technology, genomics, proteomical bioinformatics, and pharmaceutical and by medical research and techniques. "(11) SELECT AGENT OR TOXIN.—The technology identified under— "(A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	1	(B) gain of function research.
4 "(A) means the study or use of a living 5 ganism, a virus, or a product of a living or 6 nism or virus; and 7 "(B) includes each discipline, methodolog 8 and application of biology, including be 9 technology, genomics, proteomic 10 bioinformatics, and pharmaceutical and be 11 medical research and techniques. 12 "(11) SELECT AGENT OR TOXIN.—The technology identified under— 13 'select agent or toxin' means a select agent or toxidentified under— 15 "(A) section 73.3(b) of title 42, Code 16 Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; 18 "(B) section 331.3(b) of title 7, Code 19 Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; 20 or 21 "(C) section 121.3(b) of title 9, Code 23 Federal Regulations, as in effect on the date	2	"(10) Life sciences research.—The term
ganism, a virus, or a product of a living organism or virus; and "(B) includes each discipline, methodology, and application of biology, including by technology, genomics, proteomical bioinformatics, and pharmaceutical and by medical research and techniques. "(11) SELECT AGENT OR TOXIN.—The technology identified under— "(A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	3	'life sciences research'—
nism or virus; and "(B) includes each discipline, methodology and application of biology, including by technology, genomics, proteomi bioinformatics, and pharmaceutical and by medical research and techniques. "(11) SELECT AGENT OR TOXIN.—The te "(Select agent or toxin' means a select agent or toxidentified under— "(A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; or "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	4	"(A) means the study or use of a living or-
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medical research and techniques. "(11) SELECT AGENT OR TOXIN.—The te select agent or toxin' means a select agent or toxin identified under— "(A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; or "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	9	technology, genomics, proteomics,
"(11) SELECT AGENT OR TOXIN.—The te 'select agent or toxin' means a select agent or tox identified under— "(A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; "(B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; or "(C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	10	bioinformatics, and pharmaceutical and bio-
identified under— (A) section 73.3(b) of title 42, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; (B) section 331.3(b) of title 7, Code Federal Regulations, as in effect on the date enactment of the Risky Research Review Act; (B) section 331.3(b) of title 7, Code rederal Regulations, as in effect on the date enactment of the Risky Research Review Act; converges of the Risky Research Review Act; from Code and Code action 121.3(b) of title 9, Code rederal Regulations, as in effect on the date	11	medical research and techniques.
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15 "(A) section 73.3(b) of title 42, Code 16 Federal Regulations, as in effect on the date 17 enactment of the Risky Research Review Act; 18 "(B) section 331.3(b) of title 7, Code 19 Federal Regulations, as in effect on the date 20 enactment of the Risky Research Review A 21 or 22 "(C) section 121.3(b) of title 9, Code 23 Federal Regulations, as in effect on the date	13	'select agent or toxin' means a select agent or toxin
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enactment of the Risky Research Review A or (C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	18	"(B) section 331.3(b) of title 7, Code of
or (C) section 121.3(b) of title 9, Code Federal Regulations, as in effect on the date	19	Federal Regulations, as in effect on the date of
22 "(C) section 121.3(b) of title 9, Code 23 Federal Regulations, as in effect on the date	20	enactment of the Risky Research Review Act;
Federal Regulations, as in effect on the date	21	or
g ,	22	"(C) section 121.3(b) of title 9, Code of
enactment of the Risky Research Review Act.	23	Federal Regulations, as in effect on the date of
	24	enactment of the Risky Research Review Act.

1 " \S 7902. Establishment and membersh	1	"§ 7902.	Establishment	and	membershi	p
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2	"(a) Establishment.—There is established as an
3	independent agency within the Executive Branch a board
4	to be known as the 'Life Sciences Research Security
5	Board' to review proposed Federal funding for life sciences
6	research in accordance with section 7906.
7	"(b) Appointment of Members.—
8	"(1) In general.—The President shall ap-
9	point, without regard to political affiliation, 9 indi-
10	viduals who are citizens of the United States to
11	serve as members of the Board for not more than
12	2 terms of 4 years each, including—
13	"(A) the Executive Director appointed
14	under section 7903(a);
15	"(B) 5 nongovernmental scientists in a life
16	sciences field;
17	"(C) 2 nongovernmental national security
18	experts; and
19	"(D) 1 nongovernmental biosafety expert.
20	"(2) Period for nominations.—The Presi-
21	dent shall make appointments, other than the Exec-
22	utive Director, to the Board not later than 30 days
23	after the date of enactment of this chapter.
24	"(3) Considerations of Recommenda-
25	TIONS.—The President shall make appointments to
26	the Board after considering individuals rec-

1	ommended by the chair and ranking member of the
2	appropriate congressional committees.
3	"(4) Qualifications.—Individuals appointed
4	to the Board—
5	"(A) shall—
6	"(i) be impartial individuals; and
7	"(ii) be distinguished individuals of
8	high national professional reputation in
9	their respective fields who are capable of
10	exercising the independent and objective
11	judgment necessary to conduct an impar-
12	tial assessment of the potential risks and
13	benefits associated with Federal funding of
14	high-risk life sciences research to public
15	health and national security; and
16	"(B) may not be an employee on the date
17	of the appointment or during the 3-year period
18	preceding the date of the appointment.
19	"(5) Limitations.—Not more than 4 concur-
20	rent members of the Board may be an employee, a
21	subcontractor, a previous employee, or a previous
22	subcontractor of—
23	"(A) the Department of Defense;
24	"(B) the Department of Homeland Secu-
25	rity;

1	(C) the National Institute of Allergy and
2	Infectious Diseases of the Department of
3	Health and Human Services;
4	"(D) the Office of the Director of National
5	Intelligence; or
6	"(E) the Department of Energy.
7	"(6) Consideration by the senate.—
8	"(A) In General.—Nominations for ap-
9	pointment to the Executive Director of the
10	Board shall be referred to the Committee on
11	Homeland Security and Governmental Affairs
12	of the Senate for consideration.
13	"(B) RENOMINATION.—A member of the
14	Board who is recommended to serve a second
15	term shall be nominated for appointment to the
16	Board, and such nomination shall be referred
17	pursuant to subparagraph (A).
18	"(7) VACANCY.—Not later than 30 days after
19	the date on which a vacancy on the Board occurs,
20	the vacancy shall be filled in the same manner as
21	specified for the original appointment.
22	"(8) Removal.—
23	"(A) IN GENERAL.—No member of the
24	Board shall be removed from office, other than
25	by—

1	"(i) impeachment and conviction;
2	"(ii) the action of the President for
3	inefficiency, neglect of duty, malfeasance in
4	office, physical disability, mental inca-
5	pacity, or any other condition that sub-
6	stantially impairs the performance of the
7	member's duties; or
8	"(iii) the Board in accordance with
9	subparagraph (B).
10	"(B) ACTION BY BOARD.—If the Director
11	of the Office of Government Ethics determines
12	that participation by a member of the Board in
13	high-risk life sciences research constitutes a
14	conflict of interest, the Board shall take steps
15	to mitigate or manage the conflict, which may
16	include removal.
17	"(C) Notice of Removal by Presi-
18	DENT.—
19	"(i) IN GENERAL.—In the case of the
20	removal of a member of the Board by the
21	President as described in subparagraph
22	(A)(ii), not later than 10 days after the re-
23	moval, the President shall submit to the
24	chair and ranking member of the appro-
25	priate congressional committees a report

1	specifying the facts found and the grounds
2	for removal.
3	"(ii) Publication of Report.—The
4	President shall publish in the Federal Reg-
5	ister each report submitted under clause
6	(i), except that the President may, if nec-
7	essary to protect the rights of a person
8	named in the report or to prevent undue
9	interference with any pending prosecution
10	postpone or refrain from publicly pub-
11	lishing any or all of the report until the
12	completion of such pending cases or pursu-
13	ant to privacy protection requirements in
14	law.
15	"(c) Mandatory Conflicts of Interest Re-
16	VIEW.—
17	"(1) In general.—The Board, in consultation
18	with the Director of the Office of Government Eth-
19	ics, shall—
20	"(A) not later than 180 days after the date
21	of the enactment of this chapter—
22	"(i) establish criteria to determine
23	whether there is a conflict of interest with
24	respect to any individual appointed to the
25	Board, taking into consideration require-

1	ments under Federal law relating to ethics
2	requirements for employees; and
3	"(ii) upon an appointment of a mem-
4	ber to the Board under subsection (a)(1)
5	thereafter, conduct a review of each indi-
6	vidual nominated and appointed to the
7	Board to ensure the individual does not
8	have any conflict of interest under the cri-
9	teria established pursuant to clause (i):
10	and
11	"(B) periodically thereafter, conduct a re-
12	view of each individual nominated and ap-
13	pointed to the Board to ensure the individual
14	does not have any conflict of interest under the
15	criteria established pursuant to subparagraph
16	(A)(i) during the term of service of the indi-
17	vidual.
18	"(2) Notification.—
19	"(A) In general.—Not later than 3 days
20	after the date on which the Director of the Of-
21	fice of Government Ethics becomes aware that
22	a member of the Board possesses a potential
23	conflict of interest under the criteria established
24	pursuant to paragraph (1)(A)(i), the Director
25	of the Office of Government Ethics shall notify

the chair and ranking member of the appro-1 2 priate congressional committees of the potential conflict of interest. 3 4 "(B) NOTIFICATION BY MEMBER.—Not 5 later than 30 days after the date on which a 6 member of the Board becomes aware that an-7 other member of the Board possesses a poten-8 tial conflict of interest under the criteria estab-9 lished pursuant to paragraph (1)(A)(i), the 10 member of the Board or the Executive Director 11 of the Board shall notify the chair and ranking 12 member of the appropriate congressional com-13 mittees of the potential conflict of interest. 14 "(d) SECURITY CLEARANCES.—All members of the 15 Board shall be granted all the necessary security clearances and accesses, including to relevant Presidential and 16 17 department or agency special access and compartmented access programs, in an accelerated manner, subject to the 18 19 standard procedures for granting such clearances. All 20 nominees for appointment to the Board shall qualify for 21 the necessary security clearances and accesses prior to 22 being considered for confirmation by the Committee on Homeland Security and Governmental Affairs of the Sen-24 ate.

1	(e) Participation in High-risk Life Sciences
2	Research.—
3	"(1) DISCLOSURE REQUIRED.—A member of
4	the Board shall disclose whether the member has
5	participated in or is currently participating in high-
6	risk life sciences research.
7	"(2) Conflicts of interest.—
8	"(A) IN GENERAL.—The participation in
9	high-risk life sciences research by a member of
10	the Board—
11	"(i) shall be considered a potential
12	conflict of interest; and
13	"(ii) shall be subject to scrutiny by
14	the Director of the Office of Government
15	Ethics.
16	"(B) Determination.—If the Director of
17	the Office of Government Ethics determines
18	that participation by a member of the Board in
19	high-risk life sciences research constitutes a
20	conflict of interest, the Board shall take steps
21	to mitigate or manage the conflict, which may
22	include—
23	"(i) the recusal of the affected mem-
24	ber from relevant discussions and deter-
25	minations; and

1	"(ii) removal of the affected member
2	from the Board.
3	"(f) Compensation of Members.—
4	"(1) In general.—Subject to such rules as
5	may be adopted by the Board, without regard to the
6	provisions of chapter 51 and subchapter III of chap-
7	ter 53 of title 5 relating to classification and Gen-
8	eral Schedule pay rates, a member of the Board,
9	other than the Executive Director, shall be com-
10	pensated at a rate—
11	"(A) proposed by the Executive Director
12	and approved by the Board;
13	"(B) not to exceed the rate of basic pay
14	for level II of the Executive Schedule; and
15	"(C) that is commensurate with—
16	"(i) the time a member of the Board
17	spends engaged in the performance of du-
18	ties on the Board; and
19	"(ii) necessary traveling expenses.
20	"(2) Outside employment.—Subject to terms
21	and approval determined by the Director of the Of-
22	fice of Government Ethics, a member of the Board
23	may maintain outside employment and affiliations
24	while serving on the Board.
25	"(g) Oversight.—

1	"(1) Senate.—The Committee on Homeland
2	Security and Governmental Affairs of the Senate
3	shall—
4	"(A) have continuing legislative oversight
5	jurisdiction in the Senate with respect to the of-
6	ficial conduct of the Board and agency compli-
7	ance with requirements issued by the Board
8	and
9	"(B) have access to any records provided
10	to or created by the Board.
11	"(2) House of Representatives.—The Com-
12	mittee on Oversight and Accountability of the House
13	of Representatives shall—
14	"(A) have continuing legislative oversight
15	jurisdiction in the House of Representatives
16	with respect to the official conduct of the Board
17	and agency compliance with requirements
18	issued by the Board; and
19	"(B) have access to any records provided
20	to or created by the Board.
21	"(3) Duty to cooperate.—The Board shall
22	have the duty to cooperate with the exercise of over-
23	sight jurisdiction described in this subsection.
24	"(4) SECURITY CLEARANCES.—The chair and
25	ranking member of the appropriate congressional

1	committees, and designated committee staff, shall be
2	granted all security clearances and accesses held by
3	the Board, including to relevant Presidential and de-
4	partment or agency special access and compart-
5	mented access programs.
6	"(h) Office Space.—
7	"(1) In general.—In selecting office space for
8	the Board, the Board shall exhaust options for un-
9	used office spaces owned by the Federal Government
10	as of the date of enactment of this chapter.
11	"(2) Secure office space.—
12	"(A) Requests.—In order to review or
13	discuss classified information, the Board shall
14	request an accommodation from relevant agen-
15	cies to access sensitive compartmented informa-
16	tion facilities on an as-needed basis.
17	"(B) FULFILMENT.—The head of an agen-
18	cy from which the Board requests an accommo-
19	dation under subparagraph (A) shall accommo-
20	date the request in a timely manner.
21	"§ 7903. Board personnel
22	"(a) Executive Director.—
23	"(1) Appointment.—Not later than 45 days
24	after the date of enactment of this chapter, the
25	President shall appoint, by and with the advice and

1	consent of the Senate, 1 individual who is a citizen
2	of the United States, without regard to political af-
3	filiation, to the position of Executive Director of the
4	Board for a term of 4 years.
5	"(2) Qualifications.—The individual ap-
6	pointed as Executive Director under paragraph (1)
7	shall be a private individual of integrity and impar-
8	tiality who—
9	"(A) is a distinguished scientist in a life
10	sciences field; and
11	"(B) is not, and has not been for the 3-
12	year period preceding the date of the appoint-
13	ment—
14	"(i) an employee; or
15	"(ii) a participant in high-risk life
16	sciences research supported by Federal
17	funding.
18	"(3) Security Clearances.—
19	"(A) IN GENERAL.—A candidate for Exec-
20	utive Director of the Board shall be granted all
21	security clearances and accesses held by the
22	Board, including to relevant Presidential and
23	department or agency special access and com-
24	partmented access programs in an accelerated

1	manner, subject to the standard procedures for
2	granting such clearances.
3	"(B) Qualification prior to appoint
4	MENT.—The President shall ensure that a can-
5	didate for Executive Director of the Board
6	qualifies for the security clearances and ac-
7	cesses described in subparagraph (A) prior to
8	appointment.
9	"(4) Functions.—The Executive Director of
10	the Board shall—
11	"(A) serve as principal liaison to Congress
12	and agencies;
13	"(B) serve as chair of the Board;
14	"(C) be responsible for the administration
15	and coordination of the responsibilities of the
16	Board; and
17	"(D) be responsible for the administration
18	of all official activities conducted by the Board
19	"(5) Removal.—Notwithstanding section
20	7902(b)(8), the Executive Director shall not be re-
21	moved for reasons other than for cause on the
22	grounds of inefficiency, neglect of duty, malfeasance
23	in office, physical disability, mental incapacity, or
24	any other condition that substantially impairs the

1	performance of the responsibilities of the Executive
2	Director or the staff of the Board.
3	"(6) Terms.—An Executive Director of the
4	Board shall not serve more than 2 terms.
5	"(b) Staff.—
6	"(1) In general.—Without regard to the pro-
7	visions of subchapter I of chapter 33 of title 5 gov-
8	erning appointments in the competitive service, the
9	Board may appoint not more than 25 additional per-
10	sonnel to enable the Board and the Executive Direc-
11	tor to perform the duties of the Board.
12	"(2) Qualifications.—Each individual ap-
13	pointed to the staff of the Board—
14	"(A) shall be a citizen of the United States
15	of integrity and impartiality;
16	"(B) shall have expertise in the life
17	sciences field or the national security field; and
18	"(C) may not be a participant in any fed-
19	erally funded research activity on the date of
20	the appointment or during the course of service
21	of the individual on the Board.
22	"(3) Security Clearances.—
23	"(A) IN GENERAL.—A candidate for ap-
24	pointment to the staff of the Board shall be
25	granted all security clearances and accesses

1 held by the Board, including to relevant Presi-2 dential and department or agency special access 3 and compartmented access programs, in an ac-4 celerated manner, subject to the standard procedures for granting such clearances. 6 "(B) CONDITIONAL EMPLOYMENT.— 7 "(i) IN GENERAL.—The Board may 8 offer conditional employment to a can-9 didate for a staff position of the Board 10 pending the completion of security clearance background investigations. During 11 12 the pendency of such investigations, the 13 Board shall ensure that any such employee 14 does not have access to, or responsibility 15 involving, classified or otherwise restricted 16 materials. 17 UNQUALIFIED STAFF.—If the 18 Board determines that an individual hired 19 on a conditional basis under clause (i) is 20 not eligible or otherwise does not qualify 21 for all security clearances necessary to 22 carry out the responsibilities of the posi-23 tion for which conditional employment has 24 been offered, the Board shall immediately

terminate the individual's employment.

25

1	"(4) Support from agencies.—
2	"(A) IN GENERAL.—The head of each
3	agency shall designate not less than 1 full-time
4	employee of the agency as the representative of
5	the agency to—
6	"(i) provide technical assistance to the
7	Board; and
8	"(ii) support the review process of the
9	Board with respect to the agency under
10	section 7906 in a non-voting staff capacity.
11	"(B) Prohibition.—A representative of
12	an agency designated under subparagraph (A)
13	and any employee of an agency may not directly
14	or indirectly influence in any capacity a deter-
15	mination by the Board under section 7906 with
16	respect to life sciences research funded by the
17	agency.
18	"(c) Compensation.—Subject to such rules as may
19	be adopted by the Board, without regard to the provisions
20	of title 5 governing appointments in the competitive serv-
21	ice and without regard to the provisions of chapter 51 and
22	subchapter III of chapter 53 of that title relating to classi-
23	fication and General Schedule pay rates, the Executive Di-
24	rector of the Board shall—

1	"(1) be compensated at a rate not to exceed the
2	rate of basic pay for level II of the Executive Sched-
3	ule;
4	"(2) serve the entire tenure as Executive Direc-
5	tor as 1 full-time employee; and
6	"(3) appoint and fix the compensation of such
7	other personnel as may be necessary to carry out
8	this chapter.
9	"§ 7904. Board mission and functions
10	"(a) Mission.—The mission of the Board shall be
11	to issue an independent determination as to whether an
12	agency may award Federal funding for proposed high-risk
13	life sciences research, which shall be binding upon the
14	agency.
15	"(b) Powers.—The Board shall have the authority
16	to act in a manner to carry out the mission described in
17	subsection (a), including authority to—
18	"(1) prescribe regulations to carry out the re-
19	sponsibilities of the Board;
20	"(2) establish a process for the review of Fed-
21	eral funding for high-risk life sciences research prior
22	to the award of the Federal funding, which shall be
23	binding upon an agency, including information des-
24	ignated as classified or otherwise protected from dis-
25	closure;

1	(3) direct an agency to make available to the
2	Board additional information and records, including
3	information designated as classified or otherwise
4	protected from disclosure, that the Board determines
5	are required to fulfill the functions and responsibil-
6	ities Board under this chapter;
7	"(4) review any classified research conducted or
8	funded by any agency to determine whether the re-
9	search would be considered high-risk life sciences re-
10	search; and
11	"(5) through the promulgation of regulations,
12	establish processes, policies, and procedures of the
13	Board for rendering decisions under this chapter.
14	"(c) Initial Requirements.—The Board shall—
15	"(1) not later than 180 days after the date of
16	appointment of the initial members of the Board
17	under section 7902, publish procedures in the Fed-
18	eral Register establishing the process for the review
19	by the Board under section 7906;
20	"(2) prior to the establishment of the proce-
21	dures under paragraph (1), consult with the appro-
22	priate congressional committees and heads of agen-
23	cies for purposes of developing such procedures; and

1	"(3) not later than 270 days after the date of
2	the enactment of this chapter, begin carrying out the
3	duties described in section 7906.
4	"(d) Responsiveness to Congress.—Notwith-
5	standing any other provision of law, not later than 30 days
6	after the date on which the Board receives a request for
7	information from a Member of Congress, the Board shall
8	respond to the request.
9	"(e) Congressional Briefings.—Not less fre-
10	quently than quarterly, the Board shall brief the appro-
11	priate congressional committees on the work of the Board.
12	"(f) SELECT AGENT OR TOXIN UPDATES.—
13	"(1) IN GENERAL.—Not later than 15 days
14	after the date on which the Board receives a notifi-
15	cation that a select agent or toxin has been added
16	to a list of agent or toxins under a regulation de-
17	scribed in paragraph (2), the Board shall—
18	"(A) review the select agent or toxin;
19	"(B) by majority vote of members of the
20	Board, determine whether the select agent or
21	toxin should be added into the definition of 'se-
22	lect agent or toxin' under section 7901; and
23	"(C) publish any addition determined
24	under subparagraph (B) in the Federal Reg-
25	ister.

"(2) REGULATIONS DESCRIBED.—A regulation

1

2	described in this paragraph is—
3	"(A) section 73.3(b) of title 42, Code of
4	Federal Regulations, or any successor regula-
5	tion;
6	"(B) section 331.3(b) of title 7, Code of
7	Federal Regulations, or any successor regula-
8	tion; and
9	"(C) section 121.3(b) of title 9, Code of
10	Federal Regulations, or any successor regula-
11	tion.
12	"(g) Final Determination Authority.—In any
13	dispute with an agency or entity relating to the classifica-
14	tion of life sciences research under this chapter, the Board
15	shall retain final and ultimate authority in—
16	"(1) determining whether the life sciences re-
17	search is high-risk life sciences research, dual use re-
18	search of concern involving a high-consequence
19	pathogen or gain of function research;
20	"(2) interpreting definitions in section 7901;
21	and
22	"(3) determining whether a proposed Federal
23	award for life sciences research is subject to the re-
24	view process of the Board under section 7906(a)(1).

1	"§ 7905. Agency procedures; referral to Board
2	"(a) In General.—
3	"(1) Prohibition.—The head of an agency
4	may not award Federal funding for—
5	"(A) high-risk life sciences research with-
6	out approval by the Board under section
7	7906(a)(1)(B); or
8	"(B) life sciences research if the Board, in
9	accordance with section 7906(a)(2)(A)(ii), sub-
10	mits notification to the agency under section
11	7906(a)(2)(B)(i) that Board is reviewing the
12	Federal funding for life sciences research under
13	section 7906(a) until the date on which the
14	Board makes a final determination with respect
15	to the proposed Federal funding.
16	"(2) Effective date.—Paragraph (1) shall
17	take effect on the date that is 180 days after the
18	date of enactment of this chapter.
19	"(b) High-risk Attestation; Select Agent or
20	TOXIN DISCLOSURE; CERTIFICATION.—
21	"(1) In general.—An entity seeking Federal
22	funding from an agency for life sciences research
23	shall, under the penalty of perjury—
24	"(A) attest whether—
25	"(i) the life sciences research will con-
26	stitute high-risk life sciences research; and

"(ii) the entity is performing active
research with a select agent or toxin; and
"(B) if the entity is makes a positive attes-
tation under subparagraph (A), disclose the
source of funding for all active research.
"(2) ACTIVE RESEARCH WITH SELECT AGENTS
OR TOXINS.—
"(A) IN GENERAL.—The head of an agen-
cy that receives a disclosure from an entity
under paragraph (1)(B) shall submit to the
Board the disclosure.
"(B) Board inquiries.—The Board may
contact an entity that submits a disclosure
under paragraph (1)(B) to request additional
information relating to the disclosure.
"(3) Agency certification.—
"(A) Positive attestations.—The head
of an agency making an award of Federal fund-
ing to an entity that makes a positive attesta-
tion under paragraph (1)(A)(i) shall—
"(i) submit to the Board the high-risk
life sciences proposal; and
"(ii) using the process established by
the head of the agency under paragraph
(4), certify the validity of the attestation.

1	"(B) Negative attestations.—The
2	head of an agency making an award of Federa
3	funding to an entity that makes a negative at-
4	testation under paragraph (1)(A)(i) shall—
5	"(i) review the attestation; and
6	"(ii) using the process established by
7	the head of the agency under paragraph
8	(4), certify the validity of the attestation
9	"(4) Process for review.—The head of each
10	agency that awards Federal funding for life sciences
11	research, in consultation with the Board, shall estab-
12	lish and implement a process for identifying pro-
13	posals from entities seeking Federal funding for life
14	sciences research from the agency that will con-
15	stitute high-risk life sciences research.
16	"(5) Maintenance of Records.—The head of
17	each agency shall—
18	"(A) maintain records of the certification
19	process described in paragraph (3) for each ap-
20	plication for Federal funding in accordance with
21	chapter 31 of title 44; and
22	"(B) make the records maintained under
23	subparagraph (A) available for audit and review
24	upon request by the Board.
25	"(c) Notification.—

1 "(1) IN GENERAL.—Not later than 30 days be-2 fore the date on which the head of an agency plans 3 to award Federal funding to an entity for life 4 sciences research, the head of the agency shall sub-5 mit to the Board a notification of the proposed Fed-6 eral funding. 7 "(2) Contents.—The notification of Federal 8 funding for life sciences research required under 9 paragraph (1) shall include the attestation and cer-10 tification required under subsection (b). 11 "(3) Board requests.— 12 "(A) IN GENERAL.—The Board may re-13 quest additional information from the head of 14 an agency relating to a notification submitted 15 under paragraph (1). "(B) Provision of Information.—The 16 17 head of an agency from which the Board re-18 quest additional information under subpara-19 graph (A) shall provide the information in a 20 timely manner. 21 "(d) Agency Procedures.—Not later than 180 22 days after the date on which the Board publishes the proc-23 ess of the Board in the Federal Register pursuant to section 7904(c), the head of each agency shall publish on the website of the agency prepayment and preaward proce-

1	dures of the agency with respect to Federal funding for
2	life sciences research to—
3	"(1) guarantee that—
4	"(A) all high-risk life science research pro-
5	posals are referred to the Board before the
6	award of Federal funding by the agency;
7	"(B) no Federal funding for high-risk life
8	sciences research is awarded by the agency
9	without approval by the Board; and
10	"(C) not later than 30 days before the
11	date on which the head of the agency plans to
12	award the Federal funding, the agency notifies
13	the Board of the proposal for Federal funding;
14	and
15	"(2) otherwise ensure compliance with this
16	chapter.
17	"(e) Provision of Additional Information.—
18	Upon request by the Board, the head of an agency shall
19	provide any information relating to Federal funding
20	awards for life sciences research determined necessary by
21	the Board to provide oversight of the agency.
22	"(f) Change in Circumstances During Re-
23	SEARCH.—If, during the course of life sciences research
24	in progress performed by an entity supported by Federal
25	funding from an agency, circumstances arise such that the

1	life sciences research in progress may constitute high-risk
2	life sciences research in contravention to the attestation
3	of the entity under subsection (b)(1)(A)(i)—
4	"(1) the entity shall—
5	"(A) not later than 24 hours after the
6	identification of the change in circumstance,
7	pause the life sciences research in progress; and
8	"(B) not later than 5 days after the date
9	of the identification of the change in cir-
10	cumstance, submit to the head of the agency a
11	written notification through an electronic or
12	nonelectronic communication method that—
13	"(i) notifies the head of the agency of
14	the possibility that the life sciences re-
15	search in progress may constitute high-risk
16	life sciences research; and
17	"(ii) includes a detailed description of
18	each change in circumstance that may
19	transform the life sciences research in
20	progress into high-risk life sciences re-
21	search; and
22	"(2) the head of the agency shall—
23	"(A) using the process of the agency estab-
24	lished under subsection (b)(4), determine

1	whether the life sciences research in progress
2	constitutes high-risk life sciences research;
3	"(B) if the head of the agency makes a
4	negative determination under subparagraph
5	(A), inform the entity that the entity may re-
6	sume the life sciences research in progress; and
7	"(C) if the head of the agency makes a
8	positive determination under subparagraph (A),
9	immediately submit to the Board a notification
10	of the Federal funding of high-risk life sciences
11	research in progress for review under section
12	7906(a)(1).
13	"(g) Enforcement.—
14	"(1) APPLICANT REQUIREMENTS.—If an entity
15	seeking or receiving Federal funding from an agency
16	knowingly fails to make a true attestation under
17	subsection (b)(1) or promptly notify the agency of a
18	change in circumstance in accordance with sub-
19	section (f)(1), the head of the agency shall refer the
20	entity to the appropriate entity for suspension and
21	debarment proceedings relating to the receipt of
22	Federal funding.
23	"(2) Referral to inspector general.—The
24	Board shall refer any employee of an agency respon-
25	sible for overseeing and reviewing research proposals

1	relating to Federal funding that knowingly fails to
2	comply with subsection (b)(3) to the inspector gen-
3	eral of the agency.
4	"(3) Employee discipline.—
5	"(A) IN GENERAL.—The head of an agen-
6	cy employing an employee who knowingly vio-
7	lates any provision of subsection (b)(3) (or, in
8	the case of the head of an agency who violates
9	any provision of subsection (b)(3), the Presi-
10	dent) shall impose on that employee—
11	"(i) disciplinary action in accordance
12	with chapter 75 of title 5 or an equivalent
13	procedure of the agency; and
14	"(ii) permanent revocation of any ap-
15	plicable security clearance held by the em-
16	ployee.
17	"(B) CONTRACTOR PENALTY.—In the case
18	of contractor working under a contract with an
19	agency who knowingly violates subsection
20	(b)(1), the head of the agency shall refer the
21	contractor to the appropriate entity for suspen-
22	sion and debarment proceedings relating to the
23	receipt of Federal funding.
24	"(C) Employee discipline reports.—

1	"(i) In general.—Not later than
2	360 days after the date of enactment of
3	this Act, and not less frequently than once
4	every 90 days thereafter, the head of each
5	agency shall submit to the Board and the
6	appropriate congressional committees a re-
7	port that discloses, for the period covered
8	by the report, each violation by an em-
9	ployee of the agency of subsection (b)(3).
10	"(ii) Contents.—Each report sub-
11	mitted under clause (i) shall include, with
12	respect to a violation described in that
13	clause—
14	"(I) the name and professional
15	title of each employee engaged in the
16	violation;
17	"(II) a detailed explanation of
18	the nature of the violation; and
19	"(III) the date of the violation.
20	"(iii) Publication.—Not later than
21	5 days after the date on which the Board
22	receives a report under clause (i), the
23	Board shall publish on a publicly accessible
24	and searchable website the amount of vio-

1	lations that have been committed under
2	clause (i).
3	"(h) Subaward and Subcontractor Disclo-
4	SURE.—
5	"(1) In general.—During the course of high-
6	risk life sciences research in progress performed by
7	an entity supported by Federal funding from an
8	agency, the entity shall—
9	"(A) continuously disclose to the head of
10	the agency any subcontracts or subawards made
11	or planned to be made with the Federal fund-
12	ing; and
13	"(B) obtain consent from the head of the
14	agency before awarding a subcontract or award
15	described in subparagraph (A).
16	"(2) AGENCY SUBMISSION.—Not later than 30
17	days after the date on which the head of an agency
18	receives a disclosure under paragraph (1), the head
19	of the agency shall submit to the Board the disclo-
20	sure.
21	"(3) Board inquiries.—
22	"(A) IN GENERAL.—The Board may con-
23	tact an entity that submits a disclosure under
24	paragraph (1) to request additional information
25	relating to the disclosure.

1	(B) ACCESS TO REPORTS.—During the
2	course of high-risk life sciences research in
3	progress performed by an entity supported by
4	Federal funding from an agency, upon request,
5	the Board shall have access to every annual re-
6	port of—
7	"(i) the agency;
8	"(ii) the entity performing the high-
9	risk life sciences research; and
10	"(iii) any subcontractor or sub-
11	awardee of an entity described in clause
10	(ii).
12	(11).
13	"§ 7906. Board review
13	"§ 7906. Board review
13 14	"§ 7906. Board review "(a) IN GENERAL.—
131415	"(a) In General.— "(1) High-risk life sciences research.—
13 14 15 16	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the
13 14 15 16 17	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the Board receives a notification from an agency under
13 14 15 16 17 18	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the Board receives a notification from an agency under section 7905(c) relating to proposed Federal funding
13 14 15 16 17 18 19	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the Board receives a notification from an agency under section 7905(c) relating to proposed Federal funding for life sciences research that constitutes high-risk
13 14 15 16 17 18 19 20	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the Board receives a notification from an agency under section 7905(c) relating to proposed Federal funding for life sciences research that constitutes high-risk life sciences research or the Board receives a notifi-
13 14 15 16 17 18 19 20 21	"(a) In General.— "(1) High-risk life sciences research.— Not later than 120 days after the date on which the Board receives a notification from an agency under section 7905(c) relating to proposed Federal funding for life sciences research that constitutes high-risk life sciences research or the Board receives a notification from an agency under section 7905(f)(2)(C)

1	"(A) review the proposed Federal funding
2	or high-risk life sciences research in progress;
3	"(B) by a majority vote, determine wheth-
4	er the agency may award the proposed Federal
5	funding or continue to award the Federal fund-
6	ing for the high-risk life sciences research in
7	progress; and
8	"(C) by a majority vote, determine with re-
9	spect to the high-risk life sciences research
10	funded by the proposed Federal funding or
11	Federal funding for high-risk life sciences re-
12	search in progress—
13	"(i) the minimum required biosafety
14	containment level, engineering controls,
15	and operational controls;
16	"(ii) the minimum required biosecu-
17	rity engineering controls and operational
18	controls; and
19	"(iii) the minimum required personnel
20	assurance controls.
21	"(2) Proposed life sciences research.—
22	"(A) In general.—With respect to pro-
23	posed Federal funding by an agency for life
24	sciences research, the Board may—

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1	agency before the date of enactment of this Act that
2	the Board determines may constitute high-risk life
3	sciences research, the Board may—
4	"(A) direct the agency to temporarily sus-
5	pend the Federal funding;
6	"(B) require the agency to provide com-
7	plete information on the Federal funding in
8	order for the Board to complete a review of the
9	life sciences research under paragraph (1); and
10	"(C) by a majority vote of members of the
11	Board, determine whether the agency may con-
12	tinue the Federal funding.
13	"(b) Considerations.—
14	"(1) IN GENERAL.—In making a determination
15	under subsection $(a)(1)(B)$, the Board shall con-
16	sider, with respect to the high-risk life sciences re-
17	search that will be conducted with the proposed Fed-
18	eral funding or high-risk life sciences research in
19	progress—
20	"(A) whether the research poses a threat
21	to public health;
22	"(B) whether the research poses a threat
23	to public safety;
24	"(C) whether the research has a high prob-
25	ability of producing benefits for public health;

1	"(D) whether the research poses a threat
2	to large populations of animals and plants;
3	"(E) whether the research poses a threat
4	to national security;
5	"(F) whether the research is proposed to
6	be conducted at least in part in a foreign coun-
7	try;
8	"(G) the reasonably anticipated material
9	risks of the research;
10	"(H) the reasonably anticipated informa-
11	tion risks of the research;
12	"(I) the reasonably anticipated benefits of
13	the research;
14	"(J) whether the reasonably anticipated
15	benefits of the research outweigh the reasonably
16	anticipated risks; and
17	"(K) whether the benefits of the research
18	could be obtained through procedures posing
19	lower risks.
20	"(2) Weight of factors.—The presence or
21	absence of any factor under paragraph (1) shall not
22	be decisive with respect to the determination of the
23	Board under subsection (a)(1)(B).
24	"(c) Notice Following Review and Determina-
25	TION.—

1	"(1) Agency notification.—Not later than 5
2	days after the date on which the Board makes a de-
3	termination under subsection $(a)(1)(B)$ with respect
4	to Federal funding by an agency, the Executive Di-
5	rector of the Board shall notify the head of the
6	agency of the determination.
7	"(2) Board consultation.—
8	"(A) In general.—Not later than 10
9	days after receiving a notification from the
10	Board under paragraph (1), the head of an
11	agency may request a meeting with the Board
12	to discuss the determination of the Board.
13	"(B) Board response.—The Board shall
14	schedule a meeting requested by the head of an
15	agency under subparagraph (A) in a timely
16	manner.
17	"(3) Notification to appropriate congres-
18	SIONAL COMMITTEES.—If the Board determines that
19	the head of an agency may not proceed with an
20	award of proposed Federal funding under this sec-
21	tion, the Executive Director of the Board shall no-
22	tify the appropriate congressional committees when
23	the Board notifies the head of the agency.
24	"(d) Request for Expedited Review.—

1	"(1) Definition.—In this subsection, the term
2	'emergency research' means high-risk life sciences
3	research submitted to the Board that relates to a
4	public health emergency or addresses a specific na-
5	tional security concern.
6	"(2) Request; Notification.—The head of
7	an agency seeking expedited review from the Board
8	to award Federal funding for emergency research
9	shall—
10	"(A) include a request for expedited review
11	in the notification required under section
12	7905(e); and
13	"(B) on the date of the notification de-
14	scribed in subparagraph (A), submit to the
15	Board and the appropriate congressional com-
16	mittees a notification that explains why the spe-
17	cific public health emergency or national secu-
18	rity concern necessitates expedited review under
19	this subsection.
20	"(3) Internal process.—The Board shall es-
21	tablish an internal process under which the Board
22	will give proposed emergency research expedited re-
23	view under this section.
24	"(4) Temporary emergency research.—If
25	the Board does not notify the head of an agency

1	with a determination under subsection $(a)(1)(B)$
2	with respect to proposed emergency research by the
3	15 days after the date on which the head of the
4	agency submits a request under paragraph (2)(A),
5	the head of the agency may award Federal funding
6	for the emergency research on a temporary basis.
7	"(e) Scientific Expert Panels.—
8	"(1) IN GENERAL.—The Board may establish a
9	scientific panel of nongovernmental experts to advise
10	the Board in the review by the Board of life sciences
11	research pursuant to this chapter.
12	"(2) Policies and procedures.—The Board
13	shall establish and publish in the Federal Register
14	procedures and policies relating to conflicts of inter-
15	est, recusal, expertise, and related matters before
16	the establishment of the panel described in para-
17	graph (1).
18	"(3) Prohibition.—An individual serving on
19	the panel established under paragraph (1) may not
20	advise the Board on any matter with respect to
21	which the individuals has an identified or perceived
22	conflict of interest.
23	"(4) Report.—
24	"(A) IN GENERAL.—Not later than 30
25	days after the date on which the Board estab-

I	lishes a panel established under paragraph (1),
2	the Board shall submit to the appropriate con-
3	gressional committees a report that includes the
4	names, qualifications, and any identified or per-
5	ceived conflicts of interest of individuals who
6	serve on the panel.
7	"(B) Panel Changes.—Upon a change of
8	personnel on the panel established under para-
9	graph (1), the Board shall immediately submit
10	to the appropriate congressional committees an
11	update to the report required under subpara-
12	graph (A).
13	"(f) Report.—
14	"(1) In general.—Not later than 360 days
15	after the date on which the Board establishes the
16	panel described in subsection (e)(1), and annually
17	thereafter, the Board shall submit to the appropriate
18	congressional committees a report, which shall in-
19	clude a classified annex, summarizing, with respect
20	to each determination by the Board under this sec-
21	tion relating to high-risk life sciences research—
22	"(A) the findings of the Board;
23	"(B) the determination of the Board;
24	"(C) the name and location of the entity
25	proposing the life sciences research;

1	"(D) the name and location of any recipi-
2	ent of a subaward or subcontractor of an entity
3	proposing life sciences research and the nature
4	of the participation of such a recipient or sub-
5	contractor; and
6	"(E) an account of significant challenges
7	or problems, including procedural or substantive
8	challenges or problems, that arise during the
9	course of the work of the Board, including the
10	views of any member of the Board who wishes
11	to have those views included in the report.
12	"(2) Public Report.—On the date on which
13	the Board submits a report required under para-
14	graph (1), the Board shall make the report, other
15	than the classified annex included in the report,
16	available on a website.
17	"(g) Effective Date.—This section shall take ef-
18	fect on the date that is 270 days after the date of enact-
19	ment of this chapter.
20	"§ 7907. GAO Audits
21	"The Comptroller General of the United States shall
22	periodically audit the Board.

1 **"§ 7908. Funding**

- 2 "There is authorized to be appropriated to the Board
- 3 to carry out this chapter \$30,000,000 for each of fiscal
- 4 years 2025 through 2034.".
- 5 (b) CLERICAL AMENDMENT.—The table of sections
- 6 for subtitle V of title 31, United States Code, is amended
- 7 by adding at the end the following:

"Chapter 79—Life Sciences Research Security Board

- 8 (c) Financial Disclosure Reports of Board
- 9 Members.—Section 13103(f) of title 5, United States
- 10 Code, is amended—
- 11 (1) in paragraph (11), by striking "and" at the
- 12 end;
- 13 (2) in paragraph (12), by striking the period at
- the end and inserting "; and"; and
- 15 (3) by adding at the end the following:
- 16 "(13) a member of the Life Sciences Research
- 17 Security Board established under section 7902 of
- 18 title 31.".

[&]quot;7901. Definitions.

[&]quot;7902. Establishment and membership.

[&]quot;7903. Board personnel.

[&]quot;7904. Board mission and functions.

[&]quot;7905. Agency procedures; referral to Board.

[&]quot;7906. Board review.

[&]quot;7907. GAO Audits.

[&]quot;7908. Funding.".