

#### 116TH CONGRESS 2D SESSION

# S. 3224

To require the Secretary of Veterans Affairs to allow for the use of commercial institutional review boards for research, and for other purposes.

#### IN THE SENATE OF THE UNITED STATES

January 21, 2020

Mr. Cassidy (for himself and Mr. Tester) introduced the following bill; which was read twice and referred to the Committee on Veterans' Affairs

## A BILL

To require the Secretary of Veterans Affairs to allow for the use of commercial institutional review boards for research, 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 "VA Research Approval
- 5 Efficiency Act of 2020".
- 6 SEC. 2. USE BY DEPARTMENT OF VETERANS AFFAIRS OF
- 7 COMMERCIAL INSTITUTIONAL REVIEW
- 8 BOARDS IN SPONSORED RESEARCH TRIALS.
- 9 (a) In General.—Not later than 90 days after the
- 10 date of the enactment of this Act, the Secretary of Vet-

- 1 erans Affairs shall complete all necessary policy revisions
- 2 within the directive of the Veterans Health Administration
- 3 numbered 1200.05 and titled "Requirements for the Pro-
- 4 tection of Human Subjects in Research", to allow spon-
- 5 sored clinical research of the Department of Veterans Af-
- 6 fairs to use accredited commercial institutional review
- 7 boards to review research proposal protocols of the De-
- 8 partment.
- 9 (b) Identification of Review Boards.—Not later
- 10 than 90 days after the completion of the policy revisions
- 11 under subsection (a), the Secretary shall—
- 12 (1) identify accredited commercial institutional
- review boards for use in connection with sponsored
- 14 clinical research of the Department; and
- 15 (2) establish a process to modify existing ap-
- provals in the event that a commercial institutional
- 17 review board loses its accreditation during an ongo-
- ing clinical trial.
- 19 (c) Report.—
- 20 (1) IN GENERAL.—Not later than 90 days after
- 21 the completion of the policy revisions under sub-
- section (a), and annually thereafter, the Secretary
- shall submit to the Committee on Veterans' Affairs
- of the Senate and the Committee on Veterans' Af-
- fairs of the House of Representatives a report on all

1	approvals of institutional review boards, including
2	central institutional review boards and commercial
3	institutional review boards.
4	(2) Elements.—The report required by para-
5	graph (1) shall include, at a minimum, the following:
6	(A) The name of each clinical trial with re-
7	spect to which the use of an institutional review
8	board has been approved.
9	(B) The institutional review board or insti-
10	tutional review boards used in the approval
11	process for each clinical trial.
12	(C) The amount of time between submis-
13	sion and approval.
14	SEC. 3. CREATION OF OFFICE OF RESEARCH REVIEWS
14 15	SEC. 3. CREATION OF OFFICE OF RESEARCH REVIEWS  WITHIN THE OFFICE OF INFORMATION AND
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15 16 17 18	WITHIN THE OFFICE OF INFORMATION AND TECHNOLOGY OF THE DEPARTMENT OF VET- ERANS AFFAIRS.  (a) IN GENERAL.—Not later than one year after the
15 16 17 18	WITHIN THE OFFICE OF INFORMATION AND TECHNOLOGY OF THE DEPARTMENT OF VET- ERANS AFFAIRS.  (a) IN GENERAL.—Not later than one year after the date of the enactment of this Act, the Secretary of Vet-
15 16 17 18 19	WITHIN THE OFFICE OF INFORMATION AND TECHNOLOGY OF THE DEPARTMENT OF VET- ERANS AFFAIRS.  (a) IN GENERAL.—Not later than one year after the date of the enactment of this Act, the Secretary of Vet- erans Affairs shall establish within the Office of Informa-
15 16 17 18 19 20 21	WITHIN THE OFFICE OF INFORMATION AND TECHNOLOGY OF THE DEPARTMENT OF VET- ERANS AFFAIRS.  (a) IN GENERAL.—Not later than one year after the date of the enactment of this Act, the Secretary of Vet- erans Affairs shall establish within the Office of Informa- tion and Technology of the Department of Veterans Af-

1	(1) Perform centralized security reviews and
2	complete security processes for approved research
3	sponsored outside the Department, with a focus on
4	multi-site clinical trials.

- (2) Develop and maintain a list of commercially available software preferred for use in sponsored clinical trials of the Department and ensure such list is maintained as part of the official approved software products list of the Department.
- (3) Develop benchmarks for appropriate timelines for security reviews conducted by the Office.

### (c) Report.—

- (1) IN GENERAL.—Not later than one year after the establishment of the Office, the Office shall submit to the Committee on Veterans' Affairs of the Senate and the Committee on Veterans' Affairs of the House of Representatives a report on the activity of the Office.
  - (2) ELEMENTS.—The report required by paragraph (1) shall include the following:
- 22 (A) The number of security reviews completed.

1 (B) The number of personnel assigned for 2 performing the functions described in sub-3 section (b).

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