

1-1 By: Springer, Hall S.B. No. 403  
 1-2 (In the Senate - Filed January 12, 2023; February 15, 2023,  
 1-3 read first time and referred to Committee on Health & Human  
 1-4 Services; April 19, 2023, reported adversely, with favorable  
 1-5 Committee Substitute by the following vote: Yeas 9, Nays 0;  
 1-6 April 19, 2023, sent to printer.)

1-7 COMMITTEE VOTE

	Yea	Nay	Absent	PNV
1-8				
1-9	X			
1-10	X			
1-11	X			
1-12	X			
1-13	X			
1-14	X			
1-15	X			
1-16	X			
1-17	X			

1-18 COMMITTEE SUBSTITUTE FOR S.B. No. 403 By: Perry

1-19 A BILL TO BE ENTITLED  
 1-20 AN ACT

1-21 relating to a study on the adverse reactions and efficacy of  
 1-22 COVID-19 vaccines.

1-23 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF TEXAS:

1-24 SECTION 1. DEFINITIONS. In this Act:

1-25 (1) "Center" means The University of Texas Health  
 1-26 Science Center at Houston.

1-27 (2) "COVID-19" means the disease associated with  
 1-28 exposure from:

1-29 (A) the SARS-CoV-2 virus, including any variants  
 1-30 of the virus; or

1-31 (B) a COVID-19 vaccine injection.

1-32 (3) "Department" means the Department of State Health  
 1-33 Services.

1-34 (4) "Vaccine" means a substance used to stimulate  
 1-35 immunity to a particular infectious disease or pathogen.

1-36 SECTION 2. STUDY. (a) The department, in collaboration  
 1-37 with the center, shall conduct a study to assess the full scope of  
 1-38 adverse reactions, including death, and efficacy of COVID-19  
 1-39 vaccines used in this state.

1-40 (b) In conducting the study, the department shall compile  
 1-41 information regarding:

1-42 (1) the immediate short-term side effects and adverse  
 1-43 reactions experienced by vaccine recipients, including:

1-44 (A) pain, swelling, or redness at the vaccine  
 1-45 injection site;

1-46 (B) mild fever;

1-47 (C) chills;

1-48 (D) tiredness;

1-49 (E) headache; and

1-50 (F) muscle or joint aches;

1-51 (2) the immediate serious side effects and adverse  
 1-52 reactions experienced by vaccine recipients, including:

1-53 (A) difficulty breathing;

1-54 (B) swelling of the face or throat;

1-55 (C) accelerated heartbeat;

1-56 (D) body rash;

1-57 (E) dizziness; and

1-58 (F) weakness;

1-59 (3) the long-term side effects and adverse reactions  
 1-60 experienced by vaccine recipients, including:

- 2-1 (A) paralysis;
- 2-2 (B) myocarditis;
- 2-3 (C) clotting disorders;
- 2-4 (D) thrombosis with thrombocytopenia syndrome;
- 2-5 (E) Guillain-Barré syndrome;
- 2-6 (F) difficulty thinking or brain fog;
- 2-7 (G) chronic pain;
- 2-8 (H) tiredness or fatigue;
- 2-9 (I) loss of taste;
- 2-10 (J) depression;
- 2-11 (K) anxiety; and
- 2-12 (L) death; and

2-13 (4) any misrepresentation or concealment regarding  
 2-14 the efficacy or dangers of vaccination by the vaccine manufacturer  
 2-15 or a governmental agency.

2-16 (c) In conducting the study, the department and the center  
 2-17 shall compile and make available information described by  
 2-18 Subsection (b) of this section, including information collected by:

2-19 (1) the department, including information collected  
 2-20 by the department's vital statistics unit, and center;

2-21 (2) hospitals or treatment centers;

2-22 (3) any available survey;

2-23 (4) public hearings that involve health care  
 2-24 providers, researchers, injured patients, or the families of  
 2-25 injured patients speaking on COVID-19 or COVID-19 vaccine  
 2-26 experiences, including difficulties prescribing or filling  
 2-27 therapeutic prescriptions and difficulties encountered in  
 2-28 hospitals or with any medical boards; and

2-29 (5) any other relevant source, including the Vaccine  
 2-30 Adverse Event Reporting System maintained by the Centers for  
 2-31 Disease Control and Prevention and the Defense Medical Epidemiology  
 2-32 Database maintained for the United States Department of Defense.

2-33 SECTION 3. REPORT. Not later than January 1, 2024, the  
 2-34 department shall prepare and submit to the governor, lieutenant  
 2-35 governor, speaker of the house of representatives, and chairpersons  
 2-36 of legislative standing committees with jurisdiction over health  
 2-37 and safety a written report summarizing the information compiled in  
 2-38 the study and any recommendations for legislative or other action  
 2-39 to reduce the prevalence of COVID-19 vaccine side effects. The  
 2-40 department may make the report available to the public and post the  
 2-41 report on the department's Internet website.

2-42 SECTION 4. EXPIRATION. This Act expires September 1, 2025.

2-43 SECTION 5. EFFECTIVE DATE. This Act takes effect  
 2-44 immediately if it receives a vote of two-thirds of all the members  
 2-45 elected to each house, as provided by Section 39, Article III, Texas  
 2-46 Constitution. If this Act does not receive the vote necessary for  
 2-47 immediate effect, this Act takes effect September 1, 2023.

2-48 \* \* \* \* \*