HR 6133

To require the Commissioner of Food and Drugs to make available for public inspection all records of information submitted to the Food and Drug Administration in conjunction with authorizing the emergency use of, or licensing, a COVID-19 vaccine.

Congress: 117 (2021–2023, Ended)
Chamber: House
Policy Area: Health
Introduced: Dec 2, 2021
Current Status: Referred to the Subcommittee on Health.
Latest Action: Referred to the Subcommittee on Health. (Dec 3, 2021)
Official Text: https://www.congress.gov/bill/117th-congress/house-bill/6133

Sponsor

Name: Rep. Norman, Ralph [R-SC-5]
Party: Republican • **State:** SC • **Chamber:** House

Cosponsors (20 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Babin, Brian [R-TX-36]	R · TX		Dec 2, 2021
Rep. Bishop, Dan [R-NC-9]	R · NC		Dec 2, 2021
Rep. Cawthorn, Madison [R-NC-11]	R · NC		Dec 2, 2021
Rep. Duncan, Jeff [R-SC-3]	R · SC		Dec 2, 2021
Rep. Gohmert, Louie [R-TX-1]	R · TX		Dec 2, 2021
Rep. Good, Bob [R-VA-5]	R · VA		Dec 2, 2021
Rep. Gosar, Paul A. [R-AZ-4]	R · AZ		Dec 2, 2021
Rep. Greene, Marjorie Taylor [R-GA-14]	R · GA		Dec 2, 2021
Rep. Massie, Thomas [R-KY-4]	R · KY		Dec 2, 2021
Rep. Miller, Mary E. [R-IL-15]	R · IL		Dec 2, 2021
Rep. Perry, Scott [R-PA-10]	R · PA		Dec 2, 2021
Rep. Posey, Bill [R-FL-8]	R · FL		Dec 2, 2021
Rep. Roy, Chip [R-TX-21]	R · TX		Dec 2, 2021
Rep. Weber, Randy K., Sr. [R-TX-14]	R · TX		Dec 2, 2021
Rep. Webster, Daniel [R-FL-11]	R · FL		Dec 2, 2021
Rep. Cloud, Michael [R-TX-27]	R · TX		Dec 7, 2021
Rep. Harshbarger, Diana [R-TN-1]	R · TN		Dec 7, 2021
Rep. Fallon, Pat [R-TX-4]	R · TX		Dec 8, 2021
Rep. Loudermilk, Barry [R-GA-11]	R · GA		Dec 14, 2021
Rep. Franklin, C. Scott [R-FL-15]	R · FL		Mar 29, 2022

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Dec 3, 2021

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Summary (as of Dec 2, 2021)

This bill requires the Food and Drug Administration (FDA) to make publicly available all records of information submitted to the FDA in conjunction with the emergency use authorization or licensing of a COVID-19 vaccine.

Actions Timeline

- **Dec 3, 2021:** Referred to the Subcommittee on Health.
- **Dec 2, 2021:** Introduced in House
- **Dec 2, 2021:** Referred to the House Committee on Energy and Commerce.