H. R. 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.

IN THE HOUSE OF REPRESENTATIVES

DECEMBER 2, 2021

Mr. Norman (for himself, Mr. Massie, Mr. Duncan, Mr. Perry, Mr. Roy, Mr. Good of Virginia, Mr. Webster of Florida, Mrs. Miller of Illinois, Mr. Weber of Texas, Mr. Cawthorn, Mr. Posey, Mr. Bishop of North Carolina, Mr. Gohmert, Mr. Gosar, Mr. Babin, and Mrs. Greene of Georgia) introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

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.

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,
SECTION 1. PUBLIC DISCLOSURE OF INFORMATION SUBMITTED TO FDA FOR AUTHORIZING EMERGENCY USE OF, OR LICENSING, ANY COVID–19 VACCINE.

Not later than 100 days after the date of enactment of this Act, the Commissioner of Food and Drugs shall 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.