Your Name or Organization Name

Street Address

City, State, Zip

Email Address

Name of Employer

Address of Employer

City, State, Zip code

\_\_\_\_\_\_\_\_\_\_\_\_\_\_, 2021

Dear \_\_\_\_\_\_\_\_\_,

 It has come to our attention that you are mandating COVID-19 vaccination for employees at \_\_\_\_\_\_\_\_\_\_ (or that employees at \_\_\_\_\_\_\_\_\_are being coerced to receive COVID-19 vaccination, making employees uncomfortable and creating an untoward work environment).

 Experimental COVID-19 vaccines and mRNA treatments (“COVID-19 vaccination”) have not gone through the full Food and Drug Administration (FDA) approval process, and are authorized for Emergency Use only. According to 21 U.S.C. § 360bbb-3 “*Authorization for Medical Products for Use in Emergencies*,” medical products which have been granted Emergency Use Authorization may not be compulsory and must be voluntary only after full informed consent. Thus, mandate of experimental COVID-19 vaccination, or any pressure or coercion to consent to them, violates Federal law.

 Private businesses are liable for damages caused by the experimental injections if individuals are not given full freedom of choice without threat of consequences. It is interesting to note that also government officials who impose COVID-19 vaccination mandates can be held personally liable for violating Federal law, which requires voluntary and full informed consent to any product marketed under the Emergency Use Authorization. It is illegal for any public or private entity to mandate experimental COVID-19 vaccination.

 Based on 21 U.S.C. § 360bbb-3, \_\_\_\_\_\_\_\_\_\_\_ (name of Employer) is currently in violation of Federal Law by coercing its employees to receive experimental COVID-19 vaccination.

 You are responsible and will be held liable for any injuries and/or deaths which may result to employees of \_\_\_\_\_\_\_\_\_\_ (name of Employer) as a consequence of COVID-19 vaccination. Many COVID-19 inoculation injuries and deaths in the United States and abroad have been documented, with more than 4,500 deaths in the United States and more than 8,500 deaths in the European Union to date, according to the VAERS (<https://vaers.hhs.gov>) and EUDRA Vigilance (<https://www.ema.europa.eu/en/human-regulatory/research-development/pharmacovigilance/eudravigilance>), in the United States and Europe, respectively. According to a statement by the European Parliament, “following the administration of COVID-19 vaccines, there has been an extremely high incidence of serious adverse reactions that has never been hitherto detected for any medicinal product placed on the market.” (<https://www.europarl.europa.eu/doceo/document/E-9-2021-001384_EN.html>).

Please also note the following:

1. “The Moderna COVID-19 Vaccine is an unapproved vaccine.”

“The Moderna COVID-19 Vaccine has not undergone the same type of review as an FDA- approved or cleared product.”

“Serious and unexpected side effects may occur. The Moderna COVID-19 Vaccine is still being studied in clinical trials.”

https://www.cdc.gov/vaccines/covid-19/eua/modernatx.html

2. “The Pfizer-BioNTech COVID-19 Vaccine is an unapproved vaccine.”

“The Pfizer-BioNTech COVID-19 Vaccine has not undergone the same type of review as an FDA-approved or cleared product.”

“Serious and unexpected side effects may occur. Pfizer-BioNTech COVID-19 Vaccine is still being studied in clinical trials.”

<https://www.cdc.gov/vaccines/covid-19/eua/pfizer.html>

3. “The Janssen COVID-19 Vaccine is an unapproved vaccine.”

“The Janssen COVID-19 Vaccine has not undergone the same type of review as an FDA-approved or cleared product.”

“Serious and unexpected effects may occur. The Janssen COVID-19 Vaccine is still being studied in clinical trials.”

<https://www.cdc.gov/vaccines/covid-19/eua/janssen.html>

“CDC and FDA have recommended a pause in the use of the Janssen (Johnson & Johnson) COVID-19 vaccine” due to “a type of blood clot called “cerebral venous sinus thrombosis” (CVST)” which was “seen in combination with low levels of blood platelets (thrombocytopenia).”

https://www.cdc.gov/coronavirus/2019-ncov/vaccines/safety/JJUpdate.html

4. A fourth experimental COVID-19 vaccination is expected to file for Emergency Use Authorization (EUA) by the FDA, despite having caused a number of neurological disorders, blood clots, thrombocytopenia, and deaths in the US and other countries.

https://abcnews.go.com/Health/european-countries-pause-astrazeneca-covid-19-vaccine-rollout/story?id=76496321

 We hereby demand that you cease to coerce employees of \_\_\_\_\_\_\_\_\_\_\_ (name of Employer) to receive experimental COVID-19 vaccination. If you do not stop violating Federal law 21 U.S.C. § 360bbb-3, we will take all necessary legal steps for protecting our fellow citizens’ rights under Federal law.

 Please reply to email below as to whether you will cease, as required by law, to pressure employees at \_\_\_\_\_\_\_\_(name of Employer) to receive COVID-19 vaccination, effective immediately.

Sincerely,