

NOTIFICATION OF MY PROTECTION UNDER CALIFORNIA HEALTH AND SAFETY CODES

You are receiving this notice on _____ (date). I, _____, am notifying you of my rights and protection under California Health and Safety Codes, which has been codified in LAW, against medical mandates, experimentations, investigational EUA products, including failure to give fully informed consent on risks, unknowns, and long-term health impact, including possibly mandating or forcing said EUA products, but failing to communicate “experimental subject’s bill of rights,” with a list of the rights of a subject in a medical experiment, written in a language in which the subject is fluent.

Cal Health & Saf Code § 24176 § 24176. Performance of experiment without consent or willful failure to obtain consent; Damages and penalties; Waiver of rights; Effect

- (a) Any person who is primarily responsible for conduct of a medical experiment and who negligently allows the experiment to be conducted without a subject’s informed consent, as provided in this chapter, shall be liable to the subject in an amount not to exceed ten thousand dollars (\$10,000), as determined by the court. The minimum amount of damages awarded shall be five hundred dollars (\$500).
- (b) Any person who is primarily responsible for the conduct of a medical experiment and who willfully fails to obtain the subject’s informed consent, as provided in this chapter, shall be liable to the subject in an amount not to exceed twenty–five thousand dollars (\$25,000) as determined by the court. The minimum amount of damages awarded shall be one thousand dollars (\$1,000).
- (c) Any person who is primarily responsible for the conduct of a medical experiment and who willfully fails to obtain the subject’s informed consent, as provided in this chapter, and thereby exposes a subject to a known substantial risk of serious injury, either bodily harm or psychological harm, shall be guilty of a misdemeanor punishable by imprisonment in the county jail for a period not to exceed one year or a fine of fifty thousand dollars (\$50,000), or both.
- (d) Any representative or employee of a pharmaceutical company, who is directly responsible for contracting with another person for the conduct of a medical experiment, and who has knowledge of risks or hazards with respect to the experiment, and who willfully withholds information of the risks and hazards from the person contracting for the conduct of the medical experiment, and thereby exposes a subject to substantial risk of serious injury, either bodily harm or psychological harm, shall be guilty of a misdemeanor punishable by imprisonment in the county jail for a period not to exceed one year or a fine of fifty thousand dollars (\$50,000), or both.

§ 24171. Legislative intent

The Legislature hereby finds and declares that medical experimentation on human subjects is vital for the benefit of mankind, however such experimentation shall be undertaken with due respect to the preciousness of human life and the right of individuals to determine what is done to their own bodies.

The Legislature further finds and declares that:

- (a) The Nuremberg Code of Ethics in Medical Research was developed after the trial of Nazi war criminals for unethical use of persons in medical experiments; subsequently, the Declaration of Helsinki additionally established recommendations guiding doctors in experimentation involving human subjects.

§ 24172. “Experimental subject’s bill of rights”

As used in the chapter, “experimental subject’s bill of rights,” means a list of the rights of a subject in a **medical experiment, written in a language in which the subject is fluent**. Except as otherwise provided in [Section 24175](#), this list shall include, but not be limited to the subject’s right to:

- (a) Be informed of the nature and purpose of the experiment.
- (b) Be given an explanation of the procedures to be followed in the medical experiment, and any drug or device to be utilized.
- (c) Be given a description of any attendant discomforts and risks reasonably to be expected from the experiment.
- (d) Be given an explanation of any benefits to the subject reasonably to be expected from the experiment, if applicable.
- (e) Be given a disclosure of any appropriate alternative procedures, drugs or devices that might be advantageous to the subject, and their relative risks and benefits.
- (f) Be informed of the avenues of medical treatment, if any, available to the subject after the experiment if complications should arise.
- (g) Be given an opportunity to ask any questions concerning the experiment or the procedures involved.
- (h) Be instructed that consent to participate in the medical experiment may be withdrawn at any time and the subject may discontinue participation in the medical experiment without prejudice.
- (i) Be given a copy of the signed and dated written consent form as provided for by [section 24173](#) or [24178](#).
- (j) Be given the opportunity to decide to consent or not to consent to a medical experiment without the intervention of any element of force, fraud, deceit, duress, coercion, or undue influence on the subject’s decision.

§ 24174. “Medical experiment”

As used in this chapter, “medical experiment” means:

- (a) The severance or **penetration or damaging of tissues of a human subject** or the use of a drug or **device**, as defined in Section 109920 or 109925, electromagnetic radiation, heat or cold, or a biological substance or organism, in or upon a human subject in the practice or research of medicine in a manner not reasonably related to maintaining or improving the **health** of the subject or otherwise directly benefiting the subject.

The new investigational mRNA technology “penetrate” the tissues of a human subject and PCR testing are classified as a CLASS B medical device. The new mRNA vaccines did not pass the animal trials and are currently unlicensed products. By signing this letter, I have notified _____ (organization), of my right to protection against mandates from your organization. I am aware of my right to “decide to consent or not to consent to a medical experiment without the intervention of any element of force, fraud, deceit, duress, coercion, or undue influence on the subject’s decision. I am also protected by both Federal and State laws against discrimination. This notice is signed on behalf of myself or as guardian to a minor.

Printed Name _____ Date _____
 Signature _____
 Minor’s Name (Optional) _____
 Organization Notified _____
 Lawyer (Optional) _____

Received by _____ Organization _____ Date _____