Informed Consent:

The process of informed consent occurs when *communication* ***between a patient and physician*** results in the patient's authorization or agreement to undergo a specific medical intervention. In seeking a patient's informed consent (or the consent of the patient's surrogate if the patient lacks decision-making capacity or declines to participate in making decisions), physicians should:

1. Assess the patient's ability to understand relevant medical information and the implications of treatment alternatives and to make an independent, voluntary decision.
2. Present relevant information accurately and sensitively, in keeping with the patient's preferences for receiving medical information. The physician should include information about:
3. The diagnosis (when known)
4. The nature and purpose of recommended interventions
5. The burdens, risks, and expected benefits of all options, including forgoing treatment
6. Document the informed consent conversation and the patient's (or surrogate's) decision in the medical record in some manner. When the patient/surrogate has provided specific written consent, the consent form should be included in the record.

Informed Consent occurs between a patient and a doctor.

Informed Consent | American Medical Association (ama-assn.org)

VAERS COVID Vaccine Data

Vaccine Adverse Events Reporting System USA

www.openvaers.com/covid-data

571,830 Reports through August 06, 2021

12,791 DEATHS	51,242 HOSPITALIZATIONS
95,887 OFFICE VISITS	70,666 URGENT CARE
4,461 BELL'S PALSY	5,282 ANAPHYLAXIS
5,590 Heart Attacks	4,371 Myocarditis/ Pericarditis
13,140 Life Threatening	24,305 Severe Allergic Reaction
16,044 Permanently Disabled	1,505 Miscarriages
6,784 Shingles	2,554 Thrombocytopenia/ Low Platelet

Know Your Constitutional and Lawful Rights

Who's that knocking on my front door?

Solicit:

To appeal for something; to apply to for obtaining something; to ask earnestly; to ask for the purpose of receiving; to endeavor to obtain by asking or pleading; to entreat, implore, or importune; ***to make petition to;*** to plead for; to try to obtain; ***and through word implies a serious request, it requires no particular degree of importunity, entreaty, imploration, or supplication.*** People v. Phillips, 70 Cal. App.2d 449, 160 P.2d 872, 874 To awake or excite to action, or to invite. ***The term implies personal petition and importunity addressed to a particular individual to do some particular thing.***

Solicitation:

Asking; enticing; urgent request. Any action which the relation of the parties justifies in construing into a serious request.

Black's Law Dictionary 5th edition Pgs1248-1249

Connecticut Constitution

ARTICLE 1 SECTIONS 1-5 & 7, 9, 14 & 20

Section 1: All men when they form a social compact, are equal in rights; and no man or set of men are entitled to exclusive public emoluments or privileges from the community.

Section 2: All political power is inherent in the people, and all free governments are founded on their authority, and instituted for their benefit; and they have at all times an undeniable and indefeasible right to alter their form of government in such manner as they may think expedient.

Section 4: Every citizen may freely speak, write and publish his sentiments on all subjects, being responsible for the abuse of that liberty.

Section 5: No law shall ever be passed to curtail or restrain the liberty of speech or of the press.

Section 7: The people shall be secure in their persons, houses, papers and possessions from unreasonable searches or seizures; and no warrant to search any place, or to seize any person or things, shall issue without describing them as nearly as may be, nor without probable cause supported by oath or affirmation.

Section 9: No person shall be arrested, detained or punished, except in cases clearly warranted by law.

Section 14: The citizens have a right, in a peaceable manner, to assemble for their common good, and to apply to those invested with the powers of government, for redress of grievances, or other proper purposes, by petition, address or remonstrance.

Section 20: No person shall be denied the equal protection of the law nor be subjected to segregation or discrimination in the exercise or enjoyment of his or her civil or political rights because of religion, race, color, ancestry, national origin, sex or physical or mental disability.

NIH National Library of Medicine
National Center for Biotechnology Information

PubMed.gov

Informed consent disclosure to vaccine trial subjects *of risk of COVID-19 vaccines worsening clinical disease*

www.pubmed.ncbi.nim.nih.gov/33113270

Abstract

Aims of the study: Patient comprehension is a critical part of meeting medical ethics standards of informed consent in study designs. The aim of the study was to determine if sufficient literature exists to require clinicians to disclose the specific risk that COVID-19 vaccines could worsen disease upon exposure to challenge or circulating virus.

Methods used to conduct the study:

Published literature was reviewed to identify preclinical and clinical evidence that COVID-19 vaccines could worsen disease upon exposure to challenge or circulating virus. Clinical trial protocols for COVID-19 vaccines were reviewed to determine if risks were properly disclosed.

Results of the study:

COVID-19 vaccines designed to elicit neutralising antibodies may sensitise vaccine recipients to more severe disease than if they were not vaccinated. **Vaccines for SARS, MERS and RSV have never been approved,** and the data generated in the development and testing of these vaccines suggest a serious mechanistic concern: *that vaccines designed empirically using the traditional approach (consisting of the unmodified or minimally modified coronavirus viral spike to elicit neutralising antibodies), be they composed of protein, viral vector, DNA or RNA and irrespective of delivery method, may worsen COVID-19 disease via antibody-dependent enhancement (ADE).* **This risk is sufficiently obscured in clinical trial protocols and consent forms for ongoing COVID-19 vaccine trials that adequate patient comprehension of this risk is unlikely to occur, obviating truly informed consent by subjects in these trials.**

Conclusions drawn from the study and clinical implications: *The specific and significant COVID-19 risk of ADE should have been and should be prominently and independently disclosed to research subjects currently in vaccine trials, as well as those being recruited for the trials and future patients after vaccine approval, in order to meet the medical ethics standard of patient comprehension for informed consent.*

ADE: Occurs when the anti-bodies generated during an immune response recognize and bind to a pathogen, but are unable to prevent infection. *Instead, these antibodies act as a "Trojan Horse", allowing the pathogen to get into cells and exacerbate the immune response.*