

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

Reports from the Vaccine Adverse Events Reporting System.
Our data reflects all VAERS data including the "nondomestic" reports.
[read the VAERS disclaimer](#)

675,591 Reports
through September 09, 2021

106,184

OFFICE VISITS

77,919

URGENT CARE

58,440

HOSPITALIZATIONS

14,506

DEATHS

5,783

ANAPHYLAXIS

7,911

BELL'S PALSY

6,422

Heart Attacks

1,757

Miscarriages

18,439

Permanently Disabled

14,594

Life Threatening

2,910

Thrombocytopenia/
Low Platelet

7,810

Shingles

5,371

Myocarditis/
Pericarditis

27,336

Severe Allergic
Reaction

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

Nebraska Constitution

ARTICLE 1 SECTIONS 1-5, 7 & 26

Section 1: Statement of Rights

All persons are by nature free and independent and have certain inherent and inalienable rights; among these are life, liberty, the pursuit of happiness, and the right to keep and bear arms for security or defense of self, family, home, and others, and for lawful common defense, hunting, recreational use, and all other lawful purposes, and such rights shall not be denied or infringed by the state or any subdivision thereof. **To secure these rights, and the protection of property, governments are instituted among people, deriving their just powers from the consent of the governed.**

Section 2: Slavery Prohibited

There shall be neither slavery nor involuntary servitude in this state.

Section 3: Due Process of Law; Equal Protection

No person shall be deprived of life, liberty, or property, without due process of law, nor be denied equal protection of the laws.

Section 4: Religious Freedom

All persons have a natural and infeasible right to worship Almighty God according to the dictates of their own consciences. No person shall be compelled to attend, erect or support any place of worship against his consent, and no preference shall be given by law to any religious society, nor shall any interference with the rights of conscience be permitted. No religious test shall be required as a qualification for office, nor shall any person be incompetent to be a witness on account of his religious beliefs; but nothing herein shall be construed to dispense with oaths and affirmations. Religion, morality, and knowledge, however, being essential to good government, it shall be the duty of the Legislature to pass suitable laws to protect every religious denomination in the peaceable enjoyment of its own mode of public worship, and to encourage schools and the means of instruction.

Section 5: Freedom of Speech and Press

Every person may freely speak, write and publish on all subjects, being responsible for the abuse of that liberty; and in all trials for libel, both civil and criminal, the truth when published with good motives, and for justifiable ends, shall be a sufficient defense.



Section 7: Search and Seizure

The right of the people to be secure in their persons, houses, papers, and effects against unreasonable searches and seizures shall not be violated; and no warrant shall issue but upon probable cause, supported by oath or affirmation, and particularly describing the place to be searched, and the person or thing to be seized.

Section 26: Powers Retained by People

This enumeration of rights shall not be construed to impair or deny others, retained by the people, and all powers not herein delegated, remain with the people.



Informed consent

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

www.pubmed.ncbi.nlm.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.

