

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)

463,456 Reports Through

July 9th 2021

2,487
ANAPHYLAXIS

2,885
BELL'S PALSY

10,991
DEATHS

30,781
HOSPITALIZATIONS

82,535
OFFICE VISITS

59,402
URGENT CARE

3,906
Heart Attacks

2,466
Myocarditis/Pericarditis

8,832
Life Threatening

19,814
Severe Allergic Reaction

1,073
Miscarriages

9,274
Disabled

2,552
Thrombocytopenia/
Low Platelet

5,422
Tinnitus

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



Mississippi Constitution

ARTICLE 3 SECTIONS 5, 11, 13-15 & 32

Section 5: Government Originating in the People: All political power is vested in, and derived from, the people; all government of right originates with the people, is founded upon their will only, and is instituted solely for the good of the whole.

Section 11: Peaceful Assemblage; Right to Petition Government: The right of the people peaceably to assemble and petition the government on any subject shall never be impaired.

Section 13: Freedom of Speech and Press; Libel: The freedom of speech and of the press shall be held sacred; and in all prosecutions for libel the truth may be given in evidence, and the jury shall determine the law and the facts under the direction of the court; and if it shall appear to the jury that the matter charged as libelous is true, and was published with good motives and for justifiable ends, the party shall be acquitted.

Section 14: Due Process: No person shall be deprived of life, liberty, or property except by due process of law.

Section 15: Slavery and Involuntary Servitude Prohibited; Punishment for Crime: There shall be neither slavery nor involuntary servitude in this state, otherwise than in the punishment of crime, whereof the party shall have been duly convicted.

Section 32: Construction of Enumerated Rights: The enumeration of rights in this constitution shall not be construed to deny and impair others retained by, and inherent in, the people.



Informed consent disclosure to

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

www.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.