

[Investigational New Drug | IND](#)

- [Overview](#)
- [Pre-IND Process](#)
- [IND Study Protocol](#)
- [Preparing the Initial IND Submission](#)
- [Filing the IND](#)
- [Maintaining the IND](#)
- [Templates | Education | Useful Links](#)

An academic researcher may be required to submit an IND application to the FDA in order to study a marketed medical product in a new (i.e. unapproved) clinical indication. An investigator is always required to hold an IND to study an unmarketed (i.e. unapproved) medical product. In both cases, the products are considered "investigational" by FDA. The vast majority of INDs on file with the FDA are for noncommercial research.

This table provides links to information about submitting IND applications to the FDA. The resources apply to IND applications for both clinical **research** and clinical **treatment**.

IND Applications for Clinical Investigations (<i>Product Development</i>)	IND Application Reporting	IND Application Procedures	IND Application for Treatment
Overview	Overview	Overview	
Contents & Format	Protocol Amendments	Exemptions	Contents
Regulatory & Administrative Components	Information Amendments	Interactions with FDA	Treatment of Patient
Non-clinical Components	Safety Reports	Clinical Hold	Treatment of Patient in
Clinical Components	Annual Reports	Investigator's Responsibilities	Treatment

Please review the latest [FDA guidance](#) for determining the need for an IND. For additional information on submitting INDs for unmarketed medical products please contact the [HRPP](#).

Last updated: 25 Jul 2019

The Pre-IND Process

Review the five requirements below to determine if your study qualifies for exemption from an IND ([21 CFR 312.2](#)).

Still not sure? Start with the [IND Decision Tool](#). Pay particular attention to requirement #3. The [FDA Draft Guidance Investigational New Drug Applications \(INDs\) – Determining Whether Human Research Studies Can Be Conducted without an IND](#) provides more detail on a range of issues, including the process for consulting with FDA.

If you think a pre-IND meeting is warranted please contact the [HRPP](#) for assistance. Templates for a meeting request letter and pre-IND briefing packet are provided below. In addition, please consult with the IRB to determine whether a formal letter from FDA is required to document the waiver.

- [IND Exemption Letter](#)
- [Pre-IND Consultation Contact List](#)
- [Request for Pre-IND Meeting](#)
- [Pre-IND Briefing Packet](#)

IND Exemptions

A drug that is lawfully marketed in the United States is exempt from the requirements for an IND if all of the following apply:

- The investigation is not intended to be reported to the FDA in support of a new indication for use or any other significant change in the labeling for the drug.
- The investigation is not intended to support a significant change in the advertising for a prescription drug product.
- The investigation does not involve a change in route of administration, dosage level, or patient population, or other factor that significantly increases the risks (or decreases the acceptability of the risks) associated with use of the drug product.
- The investigation is conducted in compliance with the requirements for IRB review ([21 CFR 56](#)) and informed consent ([21 CFR 50](#)).
- The drug may not be represented as safe or effective for the purposes for which it is under investigation, nor may it be commercially distributed or sold.

For additional information about whether or not an IND is required for a cancer therapy drug you must contact the [NCI Regulatory Affairs Branch](#).

Need assistance or have regulatory questions? Please contact the [HRPP](#).

Last updated: 20 Sep 2019

The IND Study Protocol

A fully developed clinical protocol is the basis for both the IRB Application and the initial IND submission.

Review the [IND Protocol Template](#) for required content. A completed protocol must be included in the IND application.

Start with a *Protocol Synopsis* (page 7 and 8 of the protocol template). The protocol synopsis will be valuable if you are planning a pre-IND meeting. Compile a reference list - include all published articles and unpublished reports or manuscripts cited.

Collect a copy of each article or report listed. For approved medications, print and read the Prescribing Information.

Product information should be integrated into the protocol. In addition, your safety plan should acknowledge known safety risks from the prescribing information and incorporate relevant safety monitoring into the protocol - or show why it is not relevant to the disease under study.

Need assistance or have regulatory questions? Please contact the [HRPP](#).

Last updated: 7 Feb 2019

Prepare the Initial IND Submission

The IND Sponsor-Investigator compiles information in three broad areas:

- **Animal Pharmacology and Toxicology Studies:** Preclinical data used to assess whether the product is reasonably safe for studies in humans. For studies of marketed drugs in new indications, this section might contain data from animal models supporting the utility of the drug in the new indication.
- **Manufacturing Information:** The composition, stability, and controls used for manufacturing the drug substance and the drug product.
 - For marketed drugs, the FDA already has this information on file in the manufacturer's Drug Master File (DMF). For unmarketed drugs, the Investigator-Sponsor can request a Letter of Authorization (LOA) from the manufacturer to cross-reference the Drug Master File or existing IND if there is one. Although it is not required, the LOA is recommended.
 - For legally marketed drugs, the information in the product label or package insert might suffice for the manufacturing information.
- **Clinical Protocols and Investigator Brochures:** A detailed clinical study protocol, and Investigator Brochure are required sections of an IND application. The Investigator Brochure is primarily needed for multi-center studies and is a summary of information needed by participating investigators to assess the safety of the investigational product. For approved medications, a copy of the Prescribing Information may be used in place of an Investigator Brochure.

The IND Sponsor-Investigator writes the IND in the format of IND Application Template. The initial IND submission should be accompanied by a cover letter, an [IND Application Form - FDA 1571](#) ([see instructions](#)) and a

ClinicalTrials.Gov [Certification of Compliance - FDA 3674](#). (All forms optimized for Safari)

The IND Sponsor-Investigator must also submit a [Statement of Investigator - FDA Form FDA 1572](#) ([see instructions](#)). This form is a formal contract with FDA to adhere to Informed Consent, IRB review, and general IND regulations.

Preparing the IND application package per FDA filing requirements (number of copies, cover colors, etc):



Last updated: 25 Jul 2019

File the Initial IND and Receive an IND Number

1. The initial IND submission should be accompanied by a cover letter, a [Form FDA 1571](#) ([see instructions](#)) and a ClinicalTrials.Gov [Certification of Compliance - FDA 3674](#). (All forms optimized for Safari)
2. All must be submitted in triplicate (an original and two photocopies):
3. The IND Sponsor-Investigator completes, signs, and dates the [Form FDA 1571](#).
4. The IND Sponsor-Investigator assembles the signed submission and makes three photocopies and one PDF of the original documents.

***Please Note:** Academic Institutions are not required to submit electronically. However, if you would like to submit electronically, please see the following contacts and guidance.

- Questions and general information regarding the preparation of submissions in electronic format may be directed to CDER at esub@fda.hhs.gov or CBER at esubprep@cber.fda.gov. Questions regarding submission of datasets to CDER may be sent to edata@fda.hhs.gov.
- Please visit the [Electronic Common Technical Document \(eCTD\)](#) web page to access a wide variety of resources and support regarding eCTD submissions.

For an initial submission, the IND Sponsor-Investigator sends the original and two photocopies to the appropriate address via overnight courier. Keep one copy of the submission packet as well as a photocopy of the courier airbill.

For a Drug:

Food and Drug Administration
Center for Drug Evaluation and Research
Central Document Room
5901-B Ammendale Rd.
Beltsville, MD. 20705-1266

For a Therapeutic Biological Product:

Food and Drug Administration
Center for Drug Evaluation and Research
Therapeutic Biological Products Document Room
5901-B Ammendale Rd.
Beltsville, MD. 20705-1266

On the delivery date, track the shipment on the courier website for confirmation of delivery. Print the delivery confirmation and file it with the copy of the submission packet, which is kept in an IND Binder.

The FDA responds to the initial submission of a new IND with a letter, acknowledging receipt of the submission and assigning the IND number. The sponsor **must wait 30 calendar days before initiating any clinical trials**. During this time, FDA has an opportunity to review the IND for safety to assure that research subjects will not be subjected to unreasonable risk. If there are no issues, the **IND generally goes into effect 30 days after the Date of Receipt** shown in this letter. It is advised that you obtain written confirmation (email or fax) that the FDA review has been completed and that the study **May Proceed**.

The IND Acknowledgement letter also provides the mailing address for all subsequent submissions to the IND.

Last updated: 23 Sep 2019

Maintain the IND

To maintain an IND, the Sponsor-Investigator has three reporting responsibilities. Each type of report is time-sensitive and has a specific structure. The first two, Protocol Amendments and Safety Reports, are submitted when needed to report updated or unforeseen circumstances. The third type, the Annual Report, is submitted every year, even when no studies are in progress under the IND. Send all submissions to the address provided in the IND Acknowledgement letter received in response to the initial submission.

IND Protocol Amendments

- [Clinical Research Billing | CRB](#)
- [Human Research Protections Program | HRPP](#)
- [Office of Coverage Analysis Administration | OCAA](#)
- [Office of Contract & Grant Administration | OCGA](#)

Once an IND is in effect, the IND Sponsor-Investigator is responsible to amend it as needed to ensure that the clinical investigations are conducted according to protocols included in the application. An [IND Protocol Amendment](#) should have a [cover letter](#) and is a submission to an existing IND notifying the FDA of one or more of the following:

- New study protocol
- Change in an existing study protocol
- New investigator
- [Transfer of IND Obligations](#)

IND Annual Reports

An [IND Annual Report](#) requires a cover letter and is a brief report of the progress of studies conducted under an IND, due annually to the FDA within 60 days of the anniversary of the date that the IND went into effect.

IND Safety Reports

An [IND Safety Report](#) is expedited, written notification to the FDA of an adverse experience associated with the use of a study drug that is both serious and unexpected. “Associated with the use of the drug” is a Code of Federal Regulations term meaning “There is a reasonable possibility that the experience may have been caused by the drug.” An IND Safety Report consists of a [MedWatch Form](#) and a [cover letter](#). It is due to the FDA within 15 calendar days of initial receipt of the [SAE Report](#).

Last updated: 30 Apr 2021



- [IND Decision Tool](#)
- [IND Exemption Letter \(example 1\)](#)
- [IND Exemption Letter \(example 2\)](#)
- [Pre-IND Briefing Packet](#)
- [Pre-IND Meeting Request](#)

IND Submissions

- [FDA IND Checklist](#)
- [FDA Form 1571 - IND Application](#) and [Instructions](#) (not optimized for chrome)
- [FDA Form 1572 - IND Investigator Statement](#)
- [FDA Form 1572 \(Box 8\) - Protocol Summary Template](#)
- [FDA Form 3455 - Investigator Financial Interest Disclosure](#)
- [FDA Form 3674 - IND Certification of Compliance](#)
- [IND Application Template](#)
- [IND Cover Letter](#)
- [Protocol Template](#)
- [Request for Orphan Drug Determination](#)
- [Single Patient IND](#)
- [Expanded Access IND Submission](#)

IND Amendments

- [IND Amendment Cover Letter](#)
- [IND Protocol Amendment Template](#)
- [IND Transfer of Obligations](#)

Annual and Final Reports

- [IND Annual Report Cover Letter](#)
- [IND Annual Report Template](#)
- [IND Final Report Cover Letter](#)
- [IND Final Report Template](#)

Safety Reports

- [IND Safety Report Cover Letter](#)
- [IND Safety Report](#) and [Instructions](#) (not optimized for chrome)
- [Safety Reporting Guidance](#)

Useful Links

- [21 CFR 312](#)
- [Biological IND Submissions](#)
- [Center for Drug Evaluation & Research Guidance Documents](#) (CDER)
- [Center for Biologics Evaluation and Research Guidance Documents](#) (CBER)
- [Content and Format of IND Applications](#)
- [Exploratory INDs \(aka Phase 0\)](#)
- [FAQs about the IND Application](#)
- [FAQs about the Pre-IND Meeting](#)
- [FDA Form 1571](#)
- [FDA Form 1572](#)
- [FDA Form 3454](#)
- [FDA Form 3455](#)
- [FDA Form 3674](#)
- [Investigational New Drug Applications](#)
- [Mobile Medical Apps](#)

Last updated: 30 Apr 2021

Last updated: 20 Mar 2019

Source URL: https://compass.ucsd.edu/investigational-new-drug-ind?qt-view__vertical_tab_section__block_2=4

Drupal.jQueryUiFilter.globalOptions('accordion');