
INSTRUCTIONS FOR FILLING OUT FORM FDA 3926 – INDIVIDUAL PATIENT EXPANDED ACCESS, INVESTIGATIONAL NEW DRUG APPLICATION (IND)

(The field numbers below correspond to the numbered boxes on the Form FDA 3926.)

Do not attempt to fill out this form after opening in your Internet Browser. To enable field fillable functionality, first save the file to your local computer, close the file and re-open the Form. Checking either box 4.a or 4.b will enable the appropriate fields to be fillable. Form should be completed electronically, i.e., not hand-written.

Field 1: PHYSICIAN'S NAME, NAME OF INSTITUTION OR CLINICAL PRACTICE, ADDRESS, AND CONTACT INFORMATION

Enter the physician's name, name of institution or clinical practice, and the physician's contact information, including the physical address, email address, telephone number, and facsimile (FAX) number.

Field 2: PATIENT'S INITIALS

Enter the patient's initials (not the full name, to preserve confidentiality). The patient need not initial the form.

Field 3: DATE OF SUBMISSION

Provide the date of the submission in the following format: mm/dd/yyyy.

Field 4: TYPE OF SUBMISSION

(4.a.) Initial Submission: If the submission is an initial (original) submission for an individual patient expanded access IND (including for emergency use), select the box provided in field 4.a., enter the physician's IND number, if previously issued by FDA, and complete only fields 5 through 8, and fields 10 and 11. Do not include commercial sponsor's IND number.

(4.b.) Follow-Up Submission: If this is a follow-up submission to an existing individual patient expanded access IND, select the box provided in field 4.b. and complete the items to the right of the checkbox in field 4.b. (Investigational Drug Name and the physician's existing IND Number), and fields 9 through 11. Do not include the commercial sponsor's IND number.

Field 5: CLINICAL INFORMATION

Provide the indication (proposed treatment use) and a brief clinical history of the patient. The clinical history includes age, gender, weight, allergies or sensitivities (general (e.g. soy) and drug specific) and other optional demographic and clinical information (e.g. race (as reported by the patient; you may choose multiple answers) and ethnicity (choose only one response)), diagnosis (e.g. a brief summary (with dates) of relevant past medical and surgical history, diagnostic procedures, current stage/severity of disease, and functional status), prior therapy, response to prior therapy (e.g. patient was treated with drug X and subsequently developed lung metastasis), and the reason for requesting the proposed treatment, including an explanation of why the patient lacks other therapeutic options (e.g. patient has failed or is intolerant to currently available therapy, or is not eligible for any clinical trials registered at ClinicalTrials.gov).

Field 6: TREATMENT INFORMATION

Provide treatment information, including the investigational drug's name and the name of the entity supplying the drug (generally the manufacturer), the applicable FDA review division (if known), and a concise statement regarding the treatment plan. This includes the planned dose, route and schedule of administration of the investigational drug (including rationale for dose), planned duration of treatment, monitoring procedures (e.g. assessment criteria/procedure(s) for monitoring and frequency), planned modifications to the treatment plan in the event of toxicity (e.g. criteria for adjusting dose if dose reduction or escalation is planned, criteria for stopping the treatment), and other relevant information (e.g. concomitant

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medication). The information should be entered within the space provided. You may choose to attach an Investigator Brochure, scientific publication(s), or other supporting documents, if needed.

Field 7: LETTER OF AUTHORIZATION (LOA), IF APPLICABLE

An LOA grants FDA the right to reference another application (IND) for information to satisfy submission requirements, such as a description of the manufacturing facility, chemistry, manufacturing and controls information, and pharmacology and toxicology information.

How to obtain an LOA: The physician is responsible for obtaining the LOA in advance from the entity that is the sponsor of the IND (e.g., commercial sponsor/drug manufacturer) being referenced. Physicians should attach the LOA to Form FDA 3926. The LOA should include the IND number for the application being referenced.

If the LOA is unavailable: In cases where it is not possible to obtain an LOA (e.g., the entity supplying the drug does not have an IND already filed with FDA), physicians should contact the applicable FDA review division (see <http://www.fda.gov/NewsEvents/PublicHealthFocus/ExpandedAccessCompassionateUse/ucm429610.htm>) to determine what other sources of information may satisfy the regulatory requirements.

For emergency individual patient expanded access INDs, the physician must submit the LOA (if applicable) and all other paperwork (including Form FDA 3926) to FDA within 15 working days of FDA's initial authorization.

Field 8: PHYSICIAN'S QUALIFICATION STATEMENT

Provide a statement of the physician's qualifications. An appropriate qualification statement includes the medical school attended, year of graduation, medical specialty, state medical license number, current employment, and job title. Alternatively, the relevant portion of the physician's curriculum vitae that includes this information (usually the first few pages) may be attached.

Field 9: CONTENTS OF SUBMISSION (FOLLOW-UP/ADDITIONAL SUBMISSIONS ONLY)

This field should only be completed for follow-up/additional submissions to an existing individual patient expanded access IND. Select the appropriate box (or boxes, if more than one apply) and attach the materials indicated for the following categories of follow-up/additional submissions (the relevant FDA regulations are provided in parentheses for additional details).

If none of the following apply to the follow-up/additional communications, use Form FDA 1571 for your submission.

- *Initial Written IND Safety Report:* A report of potential serious risks to be submitted as soon as possible but no later than 15 calendar days after the sponsor (i.e., the physician, who is considered a sponsor- investigator) determines that information qualifies for reporting, or, a report of unexpected fatal or life- threatening suspected adverse reactions, submitted no later than 7 calendar days after the sponsor's initial receipt of the information (21 CFR 312.32(c))
- *Follow-up to a Written Safety Report:* A follow-up report to an IND safety report, to be made as soon as the information is available but no later than 15 calendar days after the sponsor receives the information (21 CFR 312.32(d))
- *Annual Report:* A brief report of the progress of the investigation, submitted within 60 days of the anniversary date that the IND went into effect (21 CFR 312.33)
- *Summary of Expanded Access Use (treatment completed):* A written summary of the results of the expanded access use, including adverse effects, at the conclusion of treatment (21 CFR 312.310(c)(2))

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- *Change in Treatment Plan*: Also known as protocol amendments; a submission describing changes in the IND, including changes of investigators (21 CFR 312.30)
 - *General Correspondence*: Any other communication between the sponsor and FDA pertinent to the investigation (21 CFR 312.41)
 - *Response to FDA Request for Information*: A submission containing responses to clinical information requests (21 CFR 312.41)
 - *Response to Clinical Hold*: A submission correcting deficiencies previously cited in a FDA Clinical Hold letter (21 CFR 312.42(e))
 - *Request for Withdrawal*: A submission describing the intent to withdraw an effective IND (21 CFR 312.38)

Field 10.a: REQUEST FOR AUTHORIZATION TO USE FORM FDA 3926 FOR INDIVIDUAL PATIENT EXPANDED ACCESS

Select this box to request under 21 CFR 312.10, that FDA accept the completed Form FDA 3926 to satisfy FDA's requirements for submitting an individual patient expanded access IND.

Field 10.b: REQUEST FOR AUTHORIZATION TO USE ALTERNATIVE IRB REVIEW PROCEDURES

Select this box to request under 21 CFR 56.105, authorization to obtain concurrence by the IRB chairperson or by a designated IRB member, instead of at a convened IRB meeting, before the treatment use begins, in order to comply with FDA's requirements for IRB review and approval. When a waiver is requested in this manner, the physician does not receive notice from FDA indicating that the waiver is granted.

Field 11: CERTIFICATION STATEMENT AND SIGNATURE OF THE PHYSICIAN

The licensed physician identified in Field 8 must sign this field. By signing this field, the physician certifies that treatment will not begin until 30 days after FDA receives the completed application and all required materials unless the submitting physician receives earlier notification from FDA that the treatment may proceed. The physician agrees not to begin or continue clinical investigations covered by the IND if those studies are placed on clinical hold. The physician also certifies that informed consent will be obtained in compliance with FDA requirements (see 21 CFR part 50) and that an Institutional Review Board (IRB) will be responsible for initial and continuing review and approval of the expanded access use, consistent with applicable FDA requirements (see 21 CFR part 56). The physician also acknowledges that in the case of an emergency request, treatment may begin without prior IRB approval provided the IRB is notified of the emergency treatment within 5 working days of treatment. The physician agrees to conduct the investigation in accordance with all other applicable regulatory requirements.

Information on where and how to submit this form is available at [Expanded Access – How to Submit](#)