

	<b>Mochtar Riady Institute for Nanotechnology Ethics Committee (MRIN EC)/</b>	<b>SOP/007/2020/01.4</b> <b>Effective date:</b> <b>2 January 2020</b>
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## **1. Purpose**

This standard operating procedure is designed to describe how the Secretariat of the MRIN EC manages protocol submissions to the EC

## **2. Scope**

Protocol submissions include:

- Submission for Initial Review
- Resubmission of Protocols with Corrections
- Protocol Amendment
- Continuing Review of Approved Protocols
- Protocol Termination
- Final Report / Study Report

## **3. Responsibility**

It is the responsibility of the MRIN EC Secretariat to receive, record, distribute for review and get the submission packages approved by the EC, as well as to deliver the review results to the protocol applicants.

## **4. Flow chart**

<b><u>No.</u></b>	<b><u>Activity</u></b>	<b><u>Responsibility</u></b>
1	Receive Submitted Packages	EC Secretariat
	↓	
2	Check for submission items: <input type="checkbox"/> Initial Review Application <input type="checkbox"/> Resubmission of Protocols with Corrections <input type="checkbox"/> Protocol Amendment <input type="checkbox"/> Continuing Review of Approved Protocols <input type="checkbox"/> SAE/SUSAR <input type="checkbox"/> Protocol Termination <input type="checkbox"/> Final Report / Study Report	EC Secretariat
	↓	
3	Complete the submission process	EC Secretariat
	↓	
4	Store the received packages	EC Secretariat

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## **5. Detailed instructions**

### **5.1 Receive submitted packages**

#### **5.1.1 Initial Review Application**

- Go to 5.2.

#### **5.1.2 Resubmission of Protocols with Corrections**

- Retrieve the previous receipt form from the Secretariat's records.
- Go to step 5.2.1.2

#### **5.1.3 Protocol Amendment**

- Retrieve the previous receipt form from the Secretariat's records.
- Go to step 5.2.1.3

#### **5.1.4 Continuing Review of Approved Protocols**

- Retrieve the previous receipt form from the Secretariat's records.
- Go to step 5.2.1.4

#### **5.1.5 Protocol Termination**

- Retrieve the previous receipt form from the Secretariat's records.
- Go to step 5.2.1.5

#### **5.1.6 Final Report / Study Report**

- Retrieve the previous receipt form from the Secretariat's records.
- Go to step 5.2.1.6

### **5.2 Check for submission items**

#### **5.2.1 Get relevant forms:**

##### **5.2.1.1 Initial Review Application**

- a document receipt form, AF/01-007/2020/01.4 and
- an application form for initial review (see Annex 1 , Annex 2 and Annex 3 of SOP/010/2020/01.4).
- Go to step 5.2.2.
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 should be attached).

##### **5.2.1.2 Resubmission of Protocols with corrections**

- a document receipt form (AF/01-007/2020/01.4, and

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- a review form (AF/01-012/2017/01.3 in Annex 2 (SOP/012/2020/01.4))
- Go to step 5.2
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 should be attached).

#### **5.2.1.3 Protocol Amendments**

- a document receipt form, AF/01-007/2020/01.4, and
- a re-review report form, AF/01-013/2020/01.6
- Go to step 5.2.2
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 should be attached).

#### **5.2.1.4 Annual Continuing Reviews of Approved Protocols**

- a document receipt form, AF/01-007/2020/01.4, and
- a re-review report form, AF/01-014/2020/01.4
- Go to step 5.2.2
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 should be attached).

#### **5.2.1.5 Protocol Termination**

- a document receipt form, AF/01-007/2020/01.4, and
- a re-review report form AF/01-018/2020/01.4
- Go to step 5.2.2
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 should be attached).

#### **5.2.1.6 Final Report**

- a document receipt form, AF/01-007/2020/01.4, and
- a re-review report form AF/01-015/2020/01.4
- Go to step 5.2.2
- For e-submission, go to 5.2.4 (Filled AF/01-010/2020/01.4 or AF/02-010/2020/01.4 and/or AF/03-010/2020/01.4 or should be attached).

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### 5.2.2 Fill in the forms:

- Use the form AF/01-007/2020/01.4 and fill up with relevant information submitted by PI (AF/01-010/2020/01.4, and related to submission form)

### 5.2.3 Verify Contents of Submitted Package

- Check the applicable documents to ensure that all required forms and materials are contained within the submitted package.
- Verify contents of the protocol submitted package to include Informed Consent Form
- Check completeness of necessary information in the protocol submission of the study protocol for inclusion of the followings :
  - Title of the Protocol
  - Principal Investigator
  - Sponsor
  - Abstract
  - Type of Protocol (screening, survey, clinical trial and phase)
  - Objectives
  - Anticipated Outcome
  - Inclusion/Exclusion Criteria
  - Withdrawal or discontinuation Criteria
  - Modes of Treatment Studied
  - Methodology (synopsis of study design)
  - Analysis (methods)
  - Activity plan / Timeline
  - IND Number (if applicable)
  - Schedule and Duration of Treatment
  - Efficacy or Evaluation Criteria (Response/Outcome)
  - Safety Parameters Criteria (Toxicity)
- Check the submitted **Protocol and Related Documents** for the following contents:
  - Subjects' information sheets
  - Informed Consent Form
  - Case Record Form (CRF)
  - Study budget and budget justification
  - Agreement of the study
  - Curriculum Vitae (CV) of investigators
  - Investigators' Brochure
- See if changes made to the documents be underlined or highlighted.

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#### 5.2.4. Verify electronic documents (where applicable)

- Place the electronic computer documents (protocol summary, protocol and protocol-related documents) on the MRIN EC server or the Local Area Network at the time of submission for initial protocol review or protocol amendment packages in the following drive and folder:  
D:\Ethics Committee\protocols\short name of title
- Verify that the electronic version and the contents of the documents match the copy submitted by comparing a hard copy of the electronic document with the submitted one.
- Print out the protocol documents.
- Verify the correctness of the documents.
- Check that all pages of the documents have been included and that the submitted protocol and protocol-related documents do not have missing pages.
- Certify the printed hard copy in the same manner as the submitted document(s) with the dated signature.
  
- Assign a running number to the received protocols, applying the system of 9 digits. The first four digits indicate the last two digits of the year, followed by number of the month. The next three digits are the running number of the year followed by institution code, 01 for protocol from MRIN, 02 from Faculty of Medicine (FoM)-Universitas Pelita Harapan (UPH), 03 from Siloam Hospitals, 04 from Faculty of Nursing, Universitas Pelita Harapan, 05. Department of Biology, Universitas Pelita Harapan, 06. Others. For example 1306001-01 means the first protocol submitted in June 2013 from MRIN.
- Count for correct numbers of copies.
- Store the hard copy of the electronic document with the submitted documents.
- Use the assigned running number of the protocol as the labeled name.
- Identify clearly as the hard copy of the electronic document.

#### 5.2.5 Create a Protocol Submission File

- Get the “**Protocol Submission**” file.
- Record the name and the number of the submitted protocol.
- Record the receiving date and the name of the receiver.

#### 5.3 Complete the submission process

- Check for completeness of information.
- Notify the applicants if a package is incomplete.
- State clearly the items missing in the package.
- Fill up the related parts and the missing documents.

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- Write the receiving date on the letter and the first page of the documents.
- Initial the receiver's name on the receiving documents.
- Make a photocopy of the completed Form AF/01-007/2020/01.4.
- Return the original copy of the AF/01-007/2020/01.4 to the applicants for their records.
- Keep the copy of the document receipt form in the Protocol file.
- Attach an Initial Review Application Form (AF/01-010/2020/01.4) to the Research Protocol packages.
- Keep the copy of the submitted documents with original signatures in the "Submission" file.

#### **5.4 Store the received packages**

- Bind the packages together appropriately.
- Store the dated and initial original protocol packages on the MRIN EC submission shelf for review in FIFO sequence.

### **6. Glossary**

FIFO    First In First Out sequence

IND     Investigational New Drug

### **7. Annex**

Annex 1            AF/01-007/2020/01.4 Document Receipt Form  
Annex 2            AF/02-007/2020/01.4 Document History

### **8. Reference**

- Operational Guidelines for Ethics Committees that Review Biomedical Research, World Health Organization, 2000, Geneva.
- Surveying and Evaluating Ethical Review Practices, World Health Organization, 2002, Geneva.
- International Conference on Harmonisation (ICH) Harmonised Tripartite : Guideline for Good Clinical Practice E6(R1), Current Step 4 version dated 10 June 1996, November 2005, November 2016
- Standards and Operational Guidance for Ethics Review of Health-Related Research with Human Participants, 2011, World Health Organization,

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- Pedoman Nasional Etik Penelitian Kesehatan, Komisi Nasional Etik Penelitian Kesehatan, 2011.
- Associated SOPs: SOP/008/2020/01.5, SOP/010/2020/01.4 and SOP/011/2017/01.3



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**Annex 1**  
Form AF/01-007/2020/01.4

**Document Receipt Form**

<b>Received number:</b>			
<b>Protocol Number:</b>		<b>Submitted date:</b>	
<b>Type of Submission:</b>	<input type="checkbox"/> Initial Review	<input type="checkbox"/> <b>Final Report</b>	
	<input type="checkbox"/> Resubmission for re-review	<input type="checkbox"/> Protocol Termination	
	<input type="checkbox"/> Protocol Amendments	<input type="checkbox"/> Deviation / Non-Compliance / Violation Record	
	<input type="checkbox"/> <b>SAE/SUSAR</b>	<input type="checkbox"/> Queries, Notifications and Complaints	
	<input type="checkbox"/> Continuing Review of Approved Protocols		
<b>Protocol Title:</b>			
<b>Principal Investigator:</b>			
<b>Telephone number:</b>		<b>Fax :</b>	
<b>E-mail:</b>		<b>Preferred Contact</b>	<input type="checkbox"/> Phone <input type="checkbox"/> Fax <input type="checkbox"/> e-mail
<b>Institute:</b>			
<b>Delivery route:</b>	<input type="checkbox"/> Post	<input type="checkbox"/> E-submission	<input type="checkbox"/> in Person
<b>Documents submitted:</b>	<input type="checkbox"/> complete <input type="checkbox"/> incomplete, will submit on.....		
<b>Documents to be submitted later :</b>	<input type="checkbox"/> information for subjects	Check what documents are received later on.	
	<input type="checkbox"/> informed consent form		
	<input type="checkbox"/> case report forms (CRF)	<input type="checkbox"/> information for subjects	
	<input type="checkbox"/> study budget	<input type="checkbox"/> informed consent form	
	<input type="checkbox"/> investigator's brochure	<input type="checkbox"/> case report forms (CRF)	
	<input type="checkbox"/> others.....	<input type="checkbox"/> study budget	
		<input type="checkbox"/> investigator's brochure	
		<input type="checkbox"/> others.....	
<b>Other Documents</b>			
<b>Received by:</b>			
<b>Date received:</b>			

***Note:*** Please bring this receipt with you when contacting the MRIN EC

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**Annex 2**  
Form AF/02-007/2020.01.4

Document History

Author	Version	Date	Description
EC Members	01.0	2 January 2013	<b>Final version</b>
Ivet, Lia, Mona	01.1	11 October 2014	<ol style="list-style-type: none"> <li>1. Synchronize the topic number and SOP number</li> <li>2. Format Document History : Author, Version, Date and Description of the main change</li> <li>3. Section 5.2.3. : Protocol coding to reflect a better definition of internal (MRIN) and external (UPH and Siloam Hospitals)</li> <li>4. Annex 1 : Initial Review Submitted Package</li> </ol>
Debbie S. Retnoningrum , Yan Nuryanto, Monalisa	01.2	1 April 2017	<ol style="list-style-type: none"> <li>1. Delete Komisi Etik Penelitian Kesehatan (KEPK) on the header</li> <li>2. Item 5.2.4 : Delete the word “stamp”</li> <li>3. Item 5.3 box 6: Delete the word “stam[“ and change to “write the receiving”</li> <li>4. Item 5.3 box 10 : ,Delete the sentence “Attach the filled checklist (AF/01-007/2017/01.2) with the copy of the form AF/02-007/2017/01.2 with a staple”</li> <li>5. Delete Annex 1. Annex 2 became Annex 2, add “Final report” in the Type of Submission and add “other documents” as a new line</li> </ol>
Monalisa	01.3	2 January 2019	<ol style="list-style-type: none"> <li>1. Item 5.2.4. : to change and add institution code 04. Faculty of Nursing, UPH, 05. Department of Biology, UPH, 06. Others</li> <li>2. Annex 1&amp; Flow chart : to add SAE/SUSAR in the type of submission</li> </ol>
IS,LSH,MW,	01.4	2 January 2020	<ol style="list-style-type: none"> <li>1. 5.2.1.1 , box 2 to include annex 2 &amp; 3 of SOP 010.</li> </ol>

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GE			<ol style="list-style-type: none"> <li>2. 5.2.1.1 – 5.2.1.6, box 4 : to include Annex 2 &amp; 3 of SOP</li> <li>3. 5.2.2,box 1 : replace the word of “give” with “ use “</li> <li>4. 5.2.3 Replace the “ Verify contents of the protocol submitted package to include : <ul style="list-style-type: none"> <li>• 1. Original Application Form for Initial Review..... and Protocol related documents” with “ Verify contents of the protocol submitted package to include Informed Consent Form”</li> </ul> </li> <li>5. 5.2.5 : replace the word of “ Specific File” with “Submission File”</li> <li>6. 5.3 box 1 : delete . 5.3 box 2 -12 becomes 5.3 box 1 -11</li> <li>7. 5.3,box 10 : delete the statement of “ see annex 1”</li> <li>8. Form AF 01/007 : to include deviation... and Queries, Notifications and Complaints</li> <li>9. References : <ul style="list-style-type: none"> <li>• Delete link WHO</li> <li>• Delete FERCAP SOP</li> </ul> </li> </ol>
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