

Program for the Protection of Human Subjects

Institutional Review Boards

Mount Sinai Health System One Gustave L. Levy Place, Box 1081 New York, NY 10029-6574 T 212-824-8200 F 212-876-6789 irb@mssm.edu icahn.mssm.edu/pphs

PPHS Toolkit Updates

| Document | Date | Change |
|-------------------|------------|--|
| HRP-388 | 08.01.2023 | Added language for PI change for IDS and RSC |
| | | Added language that added personnel must complete TE before |
| | | COI can begin their review |
| HRP-452 | 08.01.2023 | Added language that added personnel must complete TE before |
| | | COI can begin their review; submit once TE is completed by all |
| | | parties; add TE# to mod summary |
| HRP-502p | 07.31.2023 | Added "Authority of Authorized Representative" to signature block |
| HRP-502 | 08.15.2023 | Translated the footer |
| (11.11.2022) (all | | |
| Spanish versions) | 00.45.0000 | |
| HRP-324 | 08.15.2023 | Added section for sponsor contact |
| HRP-061 | 08.15.2023 | Added sections 5.5-5.6 per AAHRPP review |
| HRP-511 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-512B | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-512C | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-513 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-515A | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-515B | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-517 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-519 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-527 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-876 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-877 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-879 | 08.17.2023 | Removed VA references per AAHRPP review |
| HRP-232R | 11.02.2023 | Added schedule 1 drug studies to the list of studies not eligible to |
| | | use an external IRB |
| | | Other minor administrative changes |
| HRP-230 | 11.03.2023 | Expanded questions for sites to be more specific |
| HRP-098 | 04.01.2024 | Created document |
| HRP-822 | 05.30.2024 | Updated footer with J Kucera's title; |
| | | Updated Reviewing IRB #1, related to SmartIRB Section 3.4 |
| HRP-524 | 06.04.2024 | Fixed Huron bug issue in merge fields |
| HRP-212b | 07.03.2024 | Added all questions in the table format |
| | | Added a section specifically for questions relating to individuals |
| | | that have turned 18 and/or regained capacity since their data/ |
| | | specimens were collected and stored for future use. |
| HRP-503R | 07.11.2024 | Changed format of instructional text to pink text. |



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| HRP-503E | 07.11.2024 | Added instruction to delete pink text in final document. Removed mention of Sinai Central and IF# from header. Clarified that the HRP-503R is not for record/ specimen review that require consenting. Clarified location of checklists in the RUTH library Added instructions for excluding special status patients in nonconsented research. Added Setting of Human Research, and Resources Available to Conduct research sections Added instructions for excluding special status patients in nonconsented research. Changed Specimens to Samples. Removed prospective data collection option Removed option to provide information about consenting Clarified CoC eligibility criteria Combined Data Security with Data Storage Changed format of instructional text to pink text. Added instruction to delete pink text in final document. Removed mention of Sinai Central and IF# from header. Changed the Introduction section to a Study Overview section. Added a Study Procedures section. |
|---------------|------------|---|
| HRP-503 Full | 07.11.2024 | Added instructions for excluding special status patients in non- consented research. Changed Specimens to Samples. |
| TINF-303 FUII | | Changed format of instructional text to pink text. Added instruction to delete pink text in final document. Removed mention of Sinai Central and IF# from header. Hyperlinked PPHS policies and checklists. Added instructions for excluding special status patients in nonconsented research. Elaborated on the Specimen Banking session. Clarified CoC eligibility criteria Provided examples for who a prisoner is. Added instructions for when an enrolled participant becomes vulnerable. |
| HRP-930 | 07.15.2024 | Created Guidance for R2S Initial Study submission and Mount Sinai site activation |
| HRP-450 | 07.25.2024 | Added a link to eDMS website. Added the need for only one appropriate role for all study team personnel listed on the study. |



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| | | Added information about the Admin non-FCOI role. Added GCO information about industry funded studies with FACTS |
|----------|------------|--|
| | | managed contracts submitted after January 1 2024. |
| | | Updated PI proxy information. |
| | | Added information about the primary contact role. |
| HRP-451 | 07.25.2024 | Added a link to eDMS website. |
| | | Updated PI proxy information. |
| HRP-452 | 07.25.2024 | Added a link to eDMS website. |
| | | Added the need for only one appropriate role for all study team |
| | | personnel listed on the study. |
| | | Added information about the Admin non-FCOI role. |
| | | Updated PI proxy information. |
| | | Added information about the primary contact role. |
| HRP-928 | 08.05.2024 | Removed the need for consent or a waiver of consent in certain |
| | | screening procedures. |
| | | Removed mention of how to count enrolled participants at the |
| | | time of continuing review. |
| | | Removed mention of how to report screen failures at the time of |
| | | continuing review. |
| HRP-917 | 09.09.2024 | Added types of submissions that use the "site modification" |
| 327 | | submission type. |
| | | Added clarification on the type of submissions using the "update |
| | | study details" submission type. |
| HRP-212b | 10.02.2024 | Changed title from "Progress Report for Continuing Review |
| | | Specimen or Data Banking" to "Annual Activity Report for |
| | | Continuing Review Specimen or Data Banking". |
| | | Updated header with revised title. |
| | | Added clarification on when Question #2 is applicable. |
| HRP-503R | 10.14.2024 | Added clarification to the Instruction section on when this |
| | | template can be used. |
| | | • Added further instruction regarding the review period in Section 6. |
| | | Removed reference to retrospective data and the definition of |
| | | retrospective data in Section 6. |
| | | Added instruction for when data is obtained from Mount Sinai in |
| | | Section 9. |
| | | Removed the case-series box from page 5. |
| | | Added new instructions for requesting Waiver of Alterations in |
| | | Section 12. |
| HRP-935 | 10.22.2024 | Created guidance to provide instructions for the continuing review |
| | | progress report |



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| HRP-451 | 10.22.2024 | Added reference to the HRP-935 guidance document |
|---------|------------|--|
| HRP-904 | 12.09.2024 | Added clarification that consent form does not need updating |
| | | when the study is closed to enrollment and to use good judgment |
| | | for which documents need updating. |
| | | Removed outdated reference to FCOI Investigator Form. |
| HRP-451 | 12.18.2024 | Added reference for R2S forms for continuing review (HRP-820 and HRP-812). |
| HRP-450 | 12.18.2024 | Added section for the study team to include the Flesh-Kincaid |
| | | Grade Level and the Flesch-Kincaid Reading Ease Score for the |
| | | consent summary section. |
| HRP-452 | 12.18.2024 | Added statement that submissions in Clarification Request State |
| | | for four weeks will be withdrawn by PPHS |
| | | Added instructions for using the "compare" tool in RUTH |
| | | Added instructions for assigning FCOI Ancillary Review in RUTH |
| HRP-450 | 01.15.2025 | • Revised to clarify that the grant proposal must be included win the |
| | | RUTH submission but cannot serve as the standalone protocol |
| | | Revised to clarify that either the Flesh-Kincaid Grade Level <u>or</u> the |
| | | Flesch-Kincaid Reading Ease Score will need to be included (both |
| | | are not required) |
| HRP-936 | 01.15.2025 | Created guidance to provide instructions for the use of an honest |
| | | broker. |
| HRP-902 | 01.15.2025 | Added language throughout the document to clarify what |
| | | constitutes a case report and a case series; and when IRB approval |
| | | is needed. |
| HRP-454 | 02.24.2025 | Created document. |
| HRP-935 | 03.10.2025 | Reformatted the document. No new information added. |
| HRP-451 | 03.10.2025 | Added language to clarify how to report enrollment numbers. |
| | | Added language to clarify when a GCO submission is needed. |
| | | Added instructions for selecting FCOI as an ancillary review. |
| HRP-904 | 03.10.2025 | Added instructions to also select "Study Team Member |
| | | Information" as part of the modification scope. |
| | | Added instructions to update the Local Study Team Members |
| | | section of the RUTH SmartForm to reflect changes in the study |
| | | team. |
| HRP-925 | 03.20.2025 | Update the point of contact for obtaining institutional clearance. |