

The following is a comprehensive list of revisions to the CRIC MOP Final Version 2.0, dated 20070827.

Section 3.G.2.: EBT/MSCT Guidelines	Revision of eligibility requirements regarding repeat EBT/MSCT testing
Section 3.G.3.: Follow-Up of Subcohort Participants	Clarification regarding time interval of repeat EBT/MSCT testing
Section 3.I.1.: Follow-Up Visit Guidelines	Addition of information regarding Out of Clinic Visits
Section 3.I.2.: Annual Clinic Visit – 12 Months	Addition of information regarding late Echo and EBT visits
Section 3.I.5.: Annual Clinic Visit – 48 Months	Clarification of EBT schedule relating to subcohort
Section 4.B.1.: Missed Visits	Addition of additional information regarding the completion of the missed visit CRF
Section 4.B.2.: Participant Retention	Correction of instructions regarding completion of on-time/late visit CRF
Section 4.D.: Participant Transfers	Addition of instructions regarding the responsibility for communicating with and managing transfer subjects
Section 5.A.4.: General Instructions for all CRFs	Addition of instructions regarding completion of CRFs by proxy
Section 5.A.6.: ALERT_CT, ALERT_I, ALERT_U	Addition of instructions relating to the timeliness of the completion of these CRFs
Section 5.A.6.: AMPUT	Clarification regarding the completion of this CRF
Section 5.A.6.: ANCILLRY	Clarification of instructions regarding the timeliness of the completion of this CRF
Section 5.A.6.: BP	Correction of language i.e., “not obtained” to “not measured”
Section 5.A.6.: CTECH	Inclusion of specific instructions for selecting type of CT/EBT equipment used
Section 5.A.6.: CTRANS	Addition of specifications for reducing radioactive exposure
Section 5.A.6.: ECGTRANS	Addition of instructions regarding identifiers used on ECG Reports
Section 5.A.6.: ECHOTRANS	Addition of instructions regarding fax delivery of ECHOTRANS CRF to SDCC
Section 5.A.6.: ELIG	Addition of instructions regarding completion of CRF for genetic testing
Section 5.A.6.: EVENTS	Elaboration on the purpose of the EVENTS CRF
Section 5.A.6.: GFRTRANS	Elaboration on the instructions regarding the completion of the GFRTRANS CRF
Section 5.A.6.: MEDHXUP	Correction of language in directions regarding completion
Section 5.A.6.: MISSVST	Addition of language regarding annual visit time frame
Section 5.A.6.: OFFSTVST	Addition of information relating to this new CRF
Section 5.A.6.: PHYSASSESS	Revision of instructions relating to the completion of this CRF with respect to CRIC Plus
Section 5.A.6.: PROXY	Addition of information relating to this new CRF
Section 5.A.6.: PROTRANS	Clarification regarding entry and verification of PROTRANS CRF

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Section 5.A.8.: OFFSTQA – CRIC Accuracy Check for Portable Scale	Addition of instructions for this CRF used to track the portable scale for Quality Assurance
Section 5.B.1.: Queries	Inclusion of study biostatisticians in the data query process
Section 6.A.2.: Types of Alerts	Clarification of instructions regarding completion of EVENTS CRF
Section 7.A.1.: Establishing a CRIC DMS Account	Assignment of responsibility for contacting new users with account information to the Project Manager
Section 7.K.2.: View Study Alert	Replacement of original DMS Menu Image
Section 7.K.3.: DHQ Reports	Replacement of original DMS Menu Image
Section 7.K.4.: CTRANS and ECHOTRANS Data Entry	Replacement of original DMS Menu Image
Section 7.K.5.: EBT/CT Scan Report	Replacement of original DMS Menu Image
Section 9.A.3.: Instructions for Nail Specimen Collection	Addition of instructions regarding the proper storage of nail clippers
Section 9.B.1.: Anthropometric Measures	Elaboration on instructions for repeated measures of height, weight, and waist circumference
Section 9.C.4.: Method	Addition of clarification that DMS will conduct ABI calculations
Section 9.D.: BIA Procedure	Addition of statement that BIA is to be conducted annually
Section 9.D.2.: Testing Procedure	Addition of statement that BIA is to be conducted annually
Section 9.E.1.: Overview of Blood Pressure Measurement	Clarification of instructions regarding arm measurement, cuff selection and placement, and application of BP cuff
Section 9.E.2.: Blood Pressure Measurement Step by Step	Addition of instructions regarding pulse measurement; Revision of instructions relating to Standing Pulse and BP Measurement
Section 10.C.4.: “Clean Catch” Urine Proteomics Sample Collection	Clarification of original instructions
Section 10.D.1.: Collection, Processing, and Shipping of 24 HR Urine Samples	Additional information regarding inadequate urine samples and the requirements for the collection of repeat specimens; additional instructions on participants who reach ESRD
Section 10.D.2.: Sample Processing for Urine Proteomics	Clarification regarding definition of midstream urine collection
Section 12.A.1.: Patient Education	Addition of instructions relating to educating the patient about GFR testing
Section 12.B.2.: Ordering glofil	Addition of instructions regarding ordering glofil
Section 12.B.3.: GFR Test Procedure	Addition of instructions regarding the use of saline
Appendix F: Echo Procedures	Addition of MOP relating to Echo procedures
Appendix G: EBT Procedures	Addition of MOP relating to EBT procedures
Appendix H: Out of Clinic (OOC) Visit Protocol	Addition of MOP relating to Out of Clinic Visits