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

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| 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 |