	CRENAL INSUA	Participant ID:		Participant Initials:				
	OF CRICE	Clinical Center:	Site:	Visit Number:				
	COMORT STUDY	CRF Date:		RC ID:				
		(CLINIC VISIT STAT	US				
	RC completes this form to document what type of visit occurred and what was completed at this visit.							
1.	Type of Contact:			\square_1 Clinic (in person) \square_2 Phone \square_3 Offsite (in person)				
	a. If "Clinic or (Offsite" contact, where did	the visit take place?	 ☐₁ Home ☐₂ Doctor's office or healthcare clinic ☐₃ Hospital ☐₄ Nursing home/hospice ☐₅ Dialysis unit ☐₆ Other care facility ☐ෑ No in person contact, phone only ☐ଃ CRIC Research Location ☐₃8 Other location 				
2.		th versions of the CRFs address.		□ ₁ Yes □ ₀ No				
3.	Which of the follo	wing case report forms/pro	ocesses were completed	during this visit? (Check all that apply)				
	A. Physical measures: Anthropometry (PHYASSESS) Ankle Brachial Index (PHYASSESS) Bioelectric Impedance Assessment (PHYASSESS) Blood pressure (BP) Electrocardiograph (ECGTRANS) Hand Grip Dynamometer (GRIP) Physical Performance Testing (PERFORM) Balance Testing (BALANCE) B. Specimen collection: Blood draw (LABCBC, SPECIMEN, SPEC70) Urine specimen collection for proteomics (PROTRANS, PROTRANS_R) 44 hour urine specimen collection (SPECIMEN, SPEC70) Spot urine specimen collection (SPECIMEN, SPEC70) C. Research Coordinator completed case report forms: Amputation Information (AMPUT) Ancillary Studies: Participation Information (ANCILLRY) Concomitant Medications information (CMED) Modified Min-Mental Status Exam (MMSE) Renal Replacement Therapy – Primary Survey (RRTPRIM) Renal Replacement Therapy – Dialysis Unit Data Collection (RRTHD/RRTPD) General Health Questionnaire (HEALTH) Buschke Selective Reminding Test (SRT)							
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RENAL INSUIT	Participant ID:		Participant Initials:					
CRICE	Clinical Center:	Site:	Visit Number:					
COHORT STUDY	CRF Date:		RC ID:					
	C	CLINIC VISIT STAT	ับร					
RC completes	s this form to document w	hat type of visit occu	rred and what was completed at this visit.					
C Research C	oordinator completed cas	e report forms: (Cont	inued)					
C. Research Coordinator completed case report forms: (Continued)								
1 Trails A (TRAILS_A)								
☐₁ Trails A (TRAILS_A)								
CRIC Study Re-Consent Status (CONSENTII)								
☐1 Proxy Information (PROXY)								
☐ Medical Event Questionnaire (<i>EVENTSII</i>)								
\square_1 Encryption Information (Baseline only) (ENCRP)								
☐ Fracture Questionnaire (entry into Phase III) (FRACTURE)								
☐ ₁ Fracture Follow-up Questionnaire (FRACTUP)								
D. Participant	completed case report for	ms:						
☐ Beck Depression Inventory (BDI)								
☐ ₁ Diet History Questionnaire (DHQ)								
☐ ₁ Kidney Disease and Quality of Life (<i>KDQOL</i>)								
☐ ₁ Medical History (Baseline Assessment) (MEDHXII)								
□₁ Med	☐ ₁ Medical History - Update (<i>MEDHXUPIII</i>)							
□₁ Phy	☐₁ Physical Activity Assessment (PHYACT)							
☐ ₁ Symptoms List (SXLIST)								
☐ ₁ Short Test of Functional Health Literacy in Adults (STOFHLA)								
☐₁ Lubben Social Network Scale (<i>LUBBEN</i>)								

 \square_1 Adult Access to Health Care and Utilization (*HCARE*)

☐₁ Self-Efficacy Questionnaire (*EFFICACY*)

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