Table: tblCTR3CGMRunInInitiation

Synopsis:

InvID

VisitDt

This phase will instruct the subject on how to use the study CGM device. Depending on CGM use status, the subject will be asked to attempt 24/7 use of the CGM device for up to four weeks at home.

NOTE: Subjects who are Dexcom CGM users with ≥5 days/week use over past 4 weeks may skip this phase and move directly to Visit 2- Study pump training and initiation.

CGM Run-In Initiation Visit

Visit Information Section

2. Visit Date:

PtID	1. Patient ID: CTR3
Namecode	2. Initials:
ubject CGM Training	
CGMUseStatus	1. CGM use status:
	 Subject is current Dexcom CGM user 5 d/wk for last 4 weeks
	 Subject is current non-Dexcom CGM user 5 d/wk for last 4 weeks
	 Subject is non-user of CGM or CGM user <5 d/wk for last 4 weeks
DexcomTraining	2. Subject received Dexcom CGM training per protocol
	3. Sensor Insertion:
SensorDt	a. Date of Initial Sensor Insertion: / / mm/dd/yy
ReceiverSN	b. Dexcom Receiver Serial Number:

c. Dexcom Transmitter Serial Number: __

d. Dexcom Sensor Lot Number: ____

1. Name of Investigator _____

Home Use Device Preparation

TransmitterSN

SensorLotNumber

(VISITINFORMATION_A _01)

QCMeter QCMeterDs	1. Was QC testing successful with the study blood glucose meter using two different concentrations of control solution? 1a. If No, please indicate why not:
QCKetone QCKetoneDs	2. Was QC testing successful with the study blood ketone meter using two different concentrations of control solution? 2a. If No, please indicate why not:
EmergencyKit	3. Subject was given a Study Emergency Kit?